

GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:45:19 ; Search time 1184.1 seconds  
(without alignments)  
10011.921 Million cell updates/sec

Title: US-09-550-163-1  
Perfect score: 732  
Sequence: 1 caatccagaagaatccgcgt.....atgaataaagccaattt 732

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues  
Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :  
EST:  
1: em\_estba:\*  
2: em\_esthm:\*  
3: em\_estin:\*  
4: em\_estnu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: gb\_gss:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_inv:\*  
20: em\_gss\_pln:\*  
21: em\_gss\_vrt:\*  
22: em\_gss\_fun:\*  
23: em\_gss\_mam:\*  
24: em\_gss\_mus:\*  
25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rod:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	701.8	95.9	803	12	BG208163 RST27654
2	412.2	56.3	429	9	AT654552 w4ab12.x
3	393	53.7	410	9	AT1962650 wq42e03.x
4	385.8	52.7	391	9	AT1339609 q42a07.x
5	372	50.8	372	9	AT1246239 q12904.x
c 6	301.4	41.2	350	17	AZ693989 AST-2H1B5

7	295.8	40.4	1691	11	AK008619 Mus muscu
8	271.8	37.1	470	14	D85797 Rat
9	256.6	35.1	1003	12	BG261965 602373784
10	250.8	34.3	746	12	BG221966 RST41783
11	241.4	33.0	252	14	BM783832 R-EST061
12	188	25.7	188	9	AA633404 np69h11.s
13	175.4	24.0	351	13	BG938225 1AB014A12
14	119.4	16.3	121	9	AA935321 o071909.s
c 15	113	15.4	311	10	AM869303 MR3-SN006
16	101.8	13.9	314	10	BB564873 BB564873
17	66.2	9.0	272	10	BB574249 BB574249
18	61	8.3	270	10	BB595946 BB595946
19	57.2	7.8	358	10	BE486735 174950 BA
20	53.2	7.3	716	13	B1459541 603200548
21	52.8	7.2	464	9	AA667912 vvi9f07.r
22	52.8	7.2	603	9	AT956381 u174e07.y
23	52.8	7.2	674	10	BB613272 BB613272
24	52.8	7.2	754	11	AK008938 Mus muscu
25	51.8	7.1	311	10	AM869303 MR3-SN006
c 26	51	7.0	424	13	BM389584 UT-R-CN1-
27	50.8	6.9	760	14	B0194830 UT-R-CN1-
28	50.8	6.9	869	12	BF540248 BF540248
29	45.6	6.2	986	17	CNS076KL
30	42.2	5.8	484	13	Bj059218 Bj059218
c 31	42.2	5.7	687	13	Bj098508 Bj098508
32	42	5.7	597	13	Bj038615 Bj038615
33	40.4	5.5	589	13	Bj094875 Bj094875
34	40.4	5.5	629	13	Bj095114 Bj095114
35	40.4	5.5	893	14	BQ734561 AGENCOURT
36	39.8	5.4	309	13	Bj059046 Bj059046
c 37	39.8	5.4	1101	17	CNS00H80
38	39.4	5.3	713	10	AV401298 AV401298
c 39	39	5.3	552	17	AO855211 Cp61835A
c 40	39	5.3	1024	17	CNS0058T
c 41	38.4	5.2	634	10	AM187570 BMGH1777
42	38.4	5.2	694	12	BF479357 I48-3026T
43	38.4	5.2	697	13	BM301804 MCA045A01
c 44	38.2	5.2	905	17	AL077798 Drosoph11
45	37.8	5.2	433	9	AT078277 oz29905.x

## ALIGNMENTS

RESULT 1	BG208163/c	803 bp	MRNA	linear	EST 21-APR-2001
LOCUS	RST27654	Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.			
DEFINITION	BG208163				
ACCESSION	BG208163.1	GI:13729850			
VERSION	EST.				
KEYWORDS	human.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 803)				
AUTHORS	Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry,R., Cain,S., Leventhal,C., Thornton,M., Ramachandran,R., Whittington,J., Lerner,L., Costanzo,D., McElligott,K., Boozer,S., Mays,R., Smith ,E., Veloso,N., Kika,A., Hess,J., Cothren,K., Lo,K., Offenbacher ,J., Danzig,J. and Ducar,M.				
TITLE	Creation of genome-wide protein expression libraries using random activation of gene expression				
JOURNAL	Nat. Biotechnol. 19 (5), 440-445 (2001)				
MEDLINE	21227151				
COMMENT	Contact: Scott J. Cain				
	Athersys, Inc.				
	3201 Carnegie Ave, Cleveland, OH 44115, USA				
	Tel: 216 431 9900				
	Fax: 216 361 9596				
	Email: scaine@atersys.com				
FEATURES	High quality sequence stop: 550.				
	Location/Qualifiers				

Query Match	Best Local Similarity	Score	DB	Length
Matches 720; Conservative	95.9%; 98.2%;	701.8;	DB 12;	803;
	0; Mismatches 12; Indels 1; Gaps 1.			
1	CAAATCCAGAAAAGATCGGTTTCCCTAACCTTGTGCGCTATTTATTTATTTAAATTGCA	60		
Db	CAATTTTCCAGAAAAGTTCTGTTTCCCTAACCTTGTGCGCAATTTTAAATTTAAATTGCA	702		
61	CGAGAGGAGAACATGTCTACTTTTCCAAATTCACACAGACGCTGGAGAGCTCTCCG	120		
Db	CGAGAGGAGAAACATGTCTACTTTTCCAAATTCACACAGACGCTGGAGAGCTCTCCG	642		
QY	121 -AAGATTTTATTAATATATGACAAATTGGCGCCAGAACAAACAGCTGAGCAAGAG	179		
Db	641 AAGAGTTTATTAATATATGACAAATTGGCGCCAGAACAAACAGCTGAGCAAGAG	582		
QY	180 CCGTCCAGCCAAAGTTGATGCTGAGAACTTCTACTATCTCATCTGTAACCTCATGTGCA	239		
Db	581 CCGTCCAGCCAAAGTTGATGCTGAGAACTTCTACTATCTCATCTGTAACCTCATGTGCA	522		
QY	240 TGGTTGAATGTTCCTTTTCATCATCTGTCGCCATCCTGTGTGACACTGTGAATCCAGA	299		
Db	521 TGAATGAAATGTTCCTTTTCATCATCTGTCGCCATCCTGTGTGACACTGTGAATCCAGA	462		
QY	300 GACGGGAACATCCCAATGACCCCTACCCACCATTAATTTTAAGAGACTGGCAGAAAGT	359		
Db	461 GACGGGAACATCCCAATGACCCCTACCCACCATTAATTTTAAGAGACTGGCAGAAAGT	402		
QY	360 ACAAGAGCCAAATCTTGAATCTAGAAAGATCGAAGGCCACATCATGAGACATTTGGT	419		
Db	401 ACAAGAGCCAAATCTTGAATCTAGAAAGATCGAAGGCCACATCATGAGACATTTGGT	342		
QY	420 CGGCTGGGTTCAAAATGTCCTCTATTAAGGAGAGAAAGCCACCAAGCTAACATCTGACGT	479		
Db	341 CGGCTGGGTTCAAAATGTCCTCTATTAAGGAGAGAAAGCCACCAAGCTAACATCTGACGT	282		
QY	480 CCAGACATGAAGAGATGCGACAGTCCACGAGGAGCAATTCAAATTTGATCTTGAAGAA	539		
Db	281 CCAGACATGAAGAGATGCGACAGTCCACGAGGAGCAATTCAAATTTGATCTTGAAGAA	222		
QY	540 AGTGAATCTTGTCTTGTGTTGAGAAATTTTCATGAGATTAATGTTGGTGGCCAAATTAAG	599		
Db	221 AGTGAATCTTGTCTTGTGTTGAGAAATTTTCATGAGATTAATGTTGGTGGCCAAATTAAG	162		
QY	600 ATAGATGACATTTCAATCTCAGTGAATTAATGTTGCTTGTGAGACCAATTTTGTGTG	659		
Db	161 ATAGATGACATTTCAATCTCAGTGAATTAATGTTGCTTGTGAGACCAATTTTGTGTG	102		
QY	660 AAGACCTCTTTACTTTCGCGGCAAGTAAATGTCAATTTTAATCAATTAATGATGAAGA	719		
Db	101 AAGACCTCTTTACTTTCGCGGCAAGTAAATGTCAATTTTAATCAATTAATGATGAAGA	42		
QY	720 TAAAGCCAAATTT 732			
Db	41 TAGAGCCAAATTT 29			

LOCUS	AI654552	429 bp	mRNA	linear	EST 17-DEC-1999
DEFINITION	wpb4812.x1 NCI_CGAP_GC6 Homo sapiens cDNA clone IMAGE:2308895 3'				
	similar to SR:MINK_HUMAN P1532 ISK SLOW VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN ;, mRNA sequence.				
ACCESSION	AI654552				
VERSION	AI654552.1	GI:4738531			
KEYWORDS	EST.				
ORGANISM	human.				
	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.				
REFERENCE	1 (bases 1 to 429)				
AUTHORS	NCI-CGAP <a href="http://www.ncbi.nlm.nih.gov/ncicgap">http://www.ncbi.nlm.nih.gov/ncicgap</a> .				
TITLE	National Cancer Institute, Cancer Genome Anatomy Project (CGAP),				
	Tumor Gene Index				
JOURNAL	Unpublished (1997)				
COMMENT	Contact: Robert Strausberg, Ph.D. Email: cgapbs-remail.nih.gov Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D. cDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/HLNt at: <a href="http://www.bio.linnl.gov/dbrrp/image/image.html">www.bio.linnl.gov/dbrrp/image/image.html</a> Insert length: 771 Std Error: 0.00 Seq primer: -400p from Gibco High quality sequence ntpos: 411. Location/Qualifiers 1..429 /organism="Homo sapiens" /db_xref="taxon:9606" /clone="IMAGE:2308895" /clone_lib="NCI CGAP GC6" /tissue_type="Pooled germ cell tumors" /lab_host="DH10B" /note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; Plasmid DNA from the normalized library NCI_CGAP_GC4 was prepared, and as circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonetids 1257096-1258611, 1469064-1470983, and 1475592-1476743). Library constructed by Bento Soares and M. Fatima Bonaldo."				
BASE COUNT	127 a	100 c	97 g	104 t	1 others
ORIGIN					
Query Match	56.3%	Score 412.2;	DB 9;	Length 429;	
Best Local Similarity	99.0%	Pred. No. 7.1e-106;			
Matches 414:	Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0;
QY	110	GACGCTTCCCAAGGATTATTTATTA	CTATTATATGCAATTTGGCGGCAGACACACACT	169	
Db	12	GACGCTTCCCAAGGATTATTTATTA	CTATTATATGCAATTTGGCGGCAGACACACACT	71	
QY	170	GAGCAAGAGGCCCTCCACAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGAC	229		
Db	72	GAGCAAGAGGCCCTCCACAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGAC	131		
QY	230	CTCATGGATGATTTGGAATGTCTCTTTCANCAATCGGGGCATCCCTGTGAGACACTGTG	289		
Db	132	CTCATGGATGATTTGGAATGTCTCTTTCANCAATCGGGGCATCCCTGTGAGACACTGTG	191		
QY	290	AAATCCAGAGAGCGGGAACACTCCCAATGACCCCTACACACAGTACATGTTGAGAGACTGG	349		
Db	192	AAATCCAGAGAGCGGGAACACTCCCAATGACCCCTACACACAGTACATGTTGAGAGACTGG	251		
QY	350	CAGGAAAAGTCAAGAGCCAAATCTTGATATAGAGAATGGAAGGCCACCATCCATGAG	409		
Db	252	CAGGAAAAGTCAAGAGCCAAATCTTGATATAGAGAATGGAAGGCCACCATCCATGAG	311		

OY 410 AACATTGGTGGCGGTGGTTCAAAATGTCCCTGATAGAGGAGAAAGCACAAGCTAA 469  
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 DB 312 AACATTGGTGGCGGTGGTTCAAAATGTCCCTGATAGAGGAGAAAGCACAAGCTAA 371  
 |||||||  
 OY 470 CATCTGACGTCACAGATGAAGAGATGCCATGCCACAGAGGCAATTCMAATTGTCTT 527  
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 DB 372 CATCTGCGCTCCAGACATGAAGAGATGCCATGCCACAGAGGCAATTCMAATTGTCTT 429  
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 RESULT 3  
 A1962650 410 bp mRNA linear EST 08-MAR-2000  
 LOCUS wq42e03.x1 NCI-CGAP GC6 Homo sapiens cDNA clone IMAGE:2473948 3'  
 DEFINITION similar to SW:MNK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
 CHANNEL PROTEIN ;, mRNA sequence.  
 ACCESSION A1962650  
 VERSION A1962650.1 GI:5755363  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
 REFERENCE 1 (bases 1 to 410)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael  
 R. Emmert-Buck, M.D., Ph.D.  
 CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima  
 Bonaldo, Ph.D.  
 cDNA Library Arrayed by: Greg Lennon, Ph.D.  
 DNA Sequencing by: Washington University Genome Sequencing Center  
 Clone distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LNL at:  
 www.bio.lnl.gov/bdnp/image/image.html  
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 Seq primer: -40UP from Glbco.  
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 /db\_xref="taxon:9606"  
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 /clone\_lib="NCI-CGAP GC6"  
 /tissue\_type="pooled germ cell tumors"  
 /lab\_host="DH10B"  
 /note="Vector: pT73D-Pac (Pharmacia) with a modified  
 polylinker. Site.1: Not I; Site.2: Eco RI; Plasmid DNA  
 from the normalized library NCI-CGAP GC4 was prepared, and  
 58 circles were made in vitro. Following HAP purification,  
 this DNA was used as tracer in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from a pool  
 of 5,000 clones made from the same library (clones  
 1257096-1258631, 1469064-1470983, and 1475592-1476743).  
 Subtraction by Bento Soares and M. Fatima Bonaldo."  
 BASE COUNT 120 a 93 c 93 g 104 t  
 ORIGIN  
 Query Match 53.7%; Score 393; DB 9; Length 410;  
 Best Local Similarity 100.0%; Pred. No. 1.9e-100; Indels 0; Gaps 0;  
 Matches 393; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 110 GACGCTCCGAAAGATTATTATCTATATGACAATTCGCCGACACACAACAGCT 169  
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 DB 18 GACGCTCCGAAAGATTATTATCTATATGACAATTCGCCGACACACAACAGCT 77  
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 OY 170 GAGCAAGAGGCCCTCCAGCAAGAGTGTGAGAACTTCTACTATGTCATCTGTAC 229  
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 DB 78 GAGCAAGAGGCCCTCCAGCAAGAGTGTGAGAACTTCTACTATGTCATCTGTAC 137  
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OY 230 CTCATGTGATGATTGGAAATGTTCTCTTTCATCATCGTGGCCATCCCTGAGACACTGTG 289  
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 OY 350 CAGAAAAAGTACAGAGACCCAAATTTGAATCTAGAGAAATCCGAAAGGCCACCATCCATGAG 409  
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 OY 410 AACATTGGTGGCGGTGGTTCAAAATGTCCCTGATAGAGGAGAAAGCACAAGCTAA 469  
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 DB 318 AACATTGGTGGCGGTGGTTCAAAATGTCCCTGATAGAGGAGAAAGCACAAGCTAA 377  
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 OY 470 CATCTGACGTCACAGATGAAGAGATGCCAGTG 502  
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 DB 378 CATCTGACGTCACAGATGAAGAGATGCCAGTG 410  
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 A1339609 391 bp mRNA linear EST 29-DEC-1998  
 LOCUS A1339609  
 DEFINITION gq42a07.x1 Soares.NbHMPu.S1 Homo sapiens cDNA clone IMAGE:1935156  
 3' similar to SW:MNK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
 CHANNEL PROTEIN ;, mRNA sequence.  
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 VERSION A1339609.1 GI:4076536  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
 REFERENCE 1 (bases 1 to 391)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 This clone is available royalty-free through LNL; contact the  
 IMAGE Consortium (info@image.lnl.gov) for further information.  
 Seq primer: -40UP from Glbco  
 High quality sequence stop: 380.  
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 Location/Qualifiers  
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 /lab\_host="DH10B"  
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 (Pharmacia) with a modified polylinker. Site.1: Not I;  
 Site.2: Eco RI; Equal amounts of plasmid DNA from three  
 normalized libraries (melanocyte 2NDH, pregnant uterus  
 NbHPU, and fetal heart NbHHLW) were mixed, and 58 circles  
 were made in vitro. Following HAP purification, this DNA  
 was used as tracer in a subtractive hybridization  
 reaction. The driver was PCR-amplified cDNAs from pools of  
 5,000 clones made from the same 3 libraries. The pools  
 consisted of I.M.A.G.E. clones 260232-265223,  
 340488-345479, and 484488-489479."  
 BASE COUNT 119 a 93 c 93 g 86 t  
 ORIGIN  
 Query Match 52.7%; Score 385.8; DB 9; Length 391;  
 Best Local Similarity 99.5%; Pred. No. 2.1e-98; Indels 0; Gaps 0;  
 Matches 387; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 OY 111 ACGCTCCGAAAGATTATTATCTATATGACAATTCGCCGACACACAACAGCTG 170  
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Db 3 ACGTCTCCGAGAGATTTTATTACTATGCGCAATGCGCCACACACACACACCTG 62
Oy 171 AGCAAGAGGCGCCCTCCAAAGCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 230
Db 63 AGCAAGAGGCGCCCTCCAAAGCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 122
Oy 231 TCATGTGATGATGTAATGATGTTCTCTTTCATCATCATGCGGCATCCTGTGTAGACACTGTA 290
Db 123 TCATGTGATGATGTAATGATGTTCTCTTTCATCATCATGCGGCATCCTGTGTAGACACTGTA 182
Oy 291 AATCCAGAGACGGGAACTCTCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 350
Db 183 AATCCAGAGACGGGAACTCTCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 242
Oy 351 AGCAAGAGGCGCCCTCCAAAGCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 410
Db 243 AGCAAGAGGCGCCCTCCAAAGCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 302
Oy 411 ACATGTGTGGGGCTGGGTTCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 470
Db 303 ACATGTGTGGGGCTGGGTTCCAAATGATGTGAGAACTTCTACTATGTCCTGTACC 362
Oy 471 ATCTGAGTCCAGACATGAGAGATGCCA 499
Db 363 ATCTGAGTCCAGACATGAGAGATGCCA 391

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RESULT 5
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LOCUS q129904.x1 Soares_NhMHPu.S1 Homo sapiens cDNA clone IMAGE:1857942
DEFINITION 3 similar to SW:WINK_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM
CHANNEL PROTEIN ; mRNA sequence.
ACCESSION AI246239 GI:3841636
VERSION AI246239
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1 (bases 1 to 372)
AUTHORS NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLES National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaaps-f@mail.nih.gov
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
Insert Length: 921 Std Error: 0.00
Seq primer: -400P from Gibco
High quality sequence stop: 365.
FEATURES
source location/Qualifiers
1..372
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/db_xref="taxon:9606"
/clone_lib="IMAGE:1857942"
/tissue_type="Soares_NhMHPu.S1"
/issue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/notes="Organ: mixed (see below); Vector: pT73D-Pac
(Pharmacia) with a modified polylinker; Site: 1: Not I;
Site: 2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NBHM, pregnant uterus
NBHPU, and fetal heart NBH19W) were mixed, and ss circles
were made in vitro. Following Bsp purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-485479."

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BASE COUNT 115 a 89 c 86 g 82 t
ORIGIN
Query Match 50.8%; Score 372; DB 9; Length 372;
Best Local Similarity 100.0%; Pred. No. 1.7e-94;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 118 CCCAGAGATTTTATTACTATATGAGCAATTTGGCCCGACAGACAGCTGAGCAGA 177
Db 1 CGAAGAGATTTTATTACTATATGAGCAATTTGGCCCGACAGACAGCTGAGCAGA 60
Oy 178 GGGCCCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCCTGTACC 237
Db 61 GGGCCCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCCTGTACC 120
Oy 238 GATGATGGAATGTTCTCTTTCATCATCATGCGGCATCAGGAGAGACATGTAATCCAA 297
Db 121 GATGATGGAATGTTCTCTTTCATCATCATGCGGCATCAGGAGAGACATGTAATCCAA 180
Oy 298 GAGACGGGAACTCCAAATGATGATGATGATGATGATGATGATGATGATGATGATGAT 357
Db 181 GAGACGGGAACTCCAAATGATGATGATGATGATGATGATGATGATGATGATGATGAT 240
Oy 358 GTACAGAGCCAAATTTGATATGATGATGATGATGATGATGATGATGATGATGATGAT 417
Db 241 GTACAGAGCCAAATTTGATATGATGATGATGATGATGATGATGATGATGATGATGAT 300
Oy 418 TGGCGCTGGGTTCAAAATGATGATGATGATGATGATGATGATGATGATGATGATGAT 477
Db 301 TGGCGCTGGGTTCAAAATGATGATGATGATGATGATGATGATGATGATGATGATGAT 360
Oy 478 GTCCAGACATGA 489
Db 361 GTCCAGACATGA 372

```

```

RESULT 6
AZ693989/c 350 bp DNA linear GSS 18-DEC-2000
LOCUS AZ693989
DEFINITION AST-2H1B5094 Genetrap HL-60 Human Promyelocytic Leukemia Library
Homo sapiens genomic 5', DNA sequence.
ACCESSION AZ693989
VERSION AZ693989
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE 1 (bases 1 to 350)
AUTHORS Henkel,G., Llyanage,M., Pratt,E., Huang,D., Riley,M., Bernardino,A.,
Durick,K. and Pollok,B.
TITLES Exon-trap tags from a HL-60 Genomescreen(TM) Library
JOURNAL Unpublished (2000)
COMMENT Contact: Greg Henkel
Gene Expression
Auroa Biosciences Corp.
11010 Torreyana Road, San Diego, CA 92121, USA
Tel: 8584048436
Fax: 8584046719
Email: henkelg@aurobio.com
Pools of cells were isolated from a Genomescreen(TM) library. The
library of cells was generated by retroviral integration of a gene
tagging element consisting of: 1) A promoterless beta-lactamase
proceeded by a splice acceptor as a reporter for gene expression;
2) A promoter driving neomycin resistance followed by a splice
donor to trap downstream exons. 3' RACE from neomycin gene was
performed using total RNA from isolated pools. Output was shotgun
cloned in pamp-1 and used to transform DH5-alpha competent
bacteria. 5' ends of reported sequences were immediately preceded
by splice donor from the trapping construct.
Class: exon-trapped.
FEATURES
source location/Qualifiers
1..350

```

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="IMAGE:1857942"
/tissue_type="Soares_NhMHPu.S1"
/issue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"
/lab_host="DH10B"
/notes="Organ: mixed (see below); Vector: pT73D-Pac
(Pharmacia) with a modified polylinker; Site: 1: Not I;
Site: 2: Eco RI; Equal amounts of plasmid DNA from three
normalized libraries (melanocyte 2NBHM, pregnant uterus
NBHPU, and fetal heart NBH19W) were mixed, and ss circles
were made in vitro. Following Bsp purification, this DNA
was used as tracer in a subtractive hybridization
reaction. The driver was PCR-amplified cDNAs from pools of
5,000 clones made from the same 3 libraries. The pools
consisted of I.M.A.G.E. clones 260232-265223,
340488-345479, and 484488-485479."

```







QY	361	CAGACCAAACTCTGATCTTGAAAGATCGAA--GGCCACATCCATGGAACAT-TCG	417
Db	378	CAAAATGTGATCTCGATCTGGAAAGCTCCAAAGGGCCACATTCATGGAACATGGGG	437
QY	418	TCGGCGCTGGTTCAAAATGTCCCCCTGATTAAGGGAGAAAG	458
Db	438	GGCGACGGGGTTCACAGTGTCACTCGATTATTAAGCAATG	478
RESULT 10			
LOCUS	BG221966	746 bp	mRNA linear EST 21-Apr-2001
DEFINITION	BS211783	Athersys RAGE Library Homo sapiens CDNA, mRNA sequence.	
ACCESSION	BG221966		
VERSION	1	GI:13747987	
KEYWORDS	EST.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	1	(bases 1 to 746)	
AUTHORS	Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry,R., Cain,S., Leventhal,C., Thornton,M., Ramachandran,R., Whittington,J., 'E., Veloso,N., Klinka,A., Hess,U., Cochren,K., Lo,K., Offendackner, 'J., Danzig,J. and Ducar,M. Creation of genome-wide protein expression libraries using random activation of gene expression Nat. Biotechnol. 19 (5), 440-445 (2001)		
TITLE			
JOURNAL			
MEDLINE	21227151		
COMMENT	Contact: Scott J. Cain Athersys, Inc. 3201 Carnegie Ave, Cleveland, OH 44115, USA Tel: 216 431 9900 Fax: 216 361 9596 Email: scaine@atersys.com High quality sequence stop: 547. Location/Qualifiers 1..746 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_lib="Athersys RAGE Library" /cell_line="HT1080" /note="See 'Creation of Genome-wide Protein Expression Libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is HT1080, since a random activation method was used, these sequence tags are not necessarily expressed in HT1080 under normal circumstances."		
FEATURES			
source			
BASE COUNT	241 a	138 c	123 g 243 t 1 others
ORIGIN			
Query Match	34.3%	Score 250.8;	DB 12; Length 746;
Best Local Similarity	94.9%;	Pred. No. 5.6e-60;	
Matches 280;	Conservative 0;	Mismatches 13;	Indels 2; Gaps 2;
QY	1	CAATCCAGAAAAAGATCCGTTTCTCTACACTGTTCGCTATTATTTATTTAAATTGCA	60
Db	453	CAATATCCAGAAAGATCCGTTTCTCTACACTGTTCGCTATTATTTATTTAAATTGCA	512
QY	61	GCAGAGAGGAGAGATGTCTACTTTATTCCAATTTCCACAGACAGCGTGGAAAGACGTTTCG	120
Db	513	GCAGAGAGGAGAGATGTCTACTTTATTCCAATTTCCACAGACAGCGTGGAAAGACGTTTCG	572
QY	121	AAGGATTTTATTTACTTATATGAGCAATTTGGCCGCAACACAAACAGCTGAGCAGAGGC	180
Db	573	AAGGATTTTATTTACTTATATGAGCAATTTGGCCGCAACACAAACAGCTGAGCAGAGGC	632
QY	181	CCGCCAAGCAAGTGTGATCGTGAACAATTTCTACTATATGATCTGTAAGCTCATGTGAT	240
Db	633	CCGCCAAGTGTGATCGTGAACAATTTCTACTATATGATCTGTAAGCTCATGTGAT	691

QY 241 GATGGAAATGTTCTTTCAT-CATCGTGGCCATCTGGTGAGCAGCATGTGAATC 294  
|||||  
Db 692 GATTGGAATGTTCTTTTAATAAATGGGGCCATCTGGGAGCAGCATGTGAATC 746  
|||||

RESULT 11  
BM783832 252 bp mRNA linear EST 05-MAR-2002  
LOCUS K-EST0061845 S17N258215 Homo sapiens cDNA clone S17N258215-2-A12  
DEFINITION 5', mRNA sequence.  
ACCESSION BM783832  
VERSION BM783832.1 GI:19132064  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 252)  
Klim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,  
Oh,K.J., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and  
Klim,Y.S.  
21C Frontier Korean EST Project 2001  
JOURNAL Unpublished (2002)  
COMMENT Contact: Kim YS  
Genome Research Center  
Korea Research Institute of Bioscience & Biotechnology  
52 Eoan-dong Yuseong-gu, Daejeon 305-333, South Korea  
Tel: +82-42-860-4470  
Fax: +82-42-860-4409  
Email: yongsung@email.krdb.re.kr  
Plate: 2 row: A column: 12  
High quality sequence stop: 252.  
Location/Qualifiers  
1..252  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="S17N258215-2-A12"  
/clone\_id="S17N258215"  
/sex="M"  
/lab\_host="TOP10F"  
/note="Organ: Stomach; Vector: pcns; site:1; EcoRI:  
site\_2; NotI: The poly (A)+ RNA was dephosphorylated with  
bacterial alkaline phosphatase (BAP) and then deprotected  
with tobacco acid pyrophosphatase (TAP). The deprotected  
intact mRNA was ligated with DNA-RNA linker including EcoR  
I site by treatment of T4 RNA ligase and the first strand  
cDNA was synthesized from oligo dt-selected mRNA by  
priming with dt-tailed vector. The dt-tailed vector was  
adjusted to have about 60nt. The cDNA vector was  
circularized with E. coli DNA ligase after digestion of  
EcoRI which site is also included in vector. An RNA strand  
converted to a DNA strand by Okayama-Berg method. The  
obtained cDNA vectors were used for transformation of  
competent cells E. coli TOP10F by electroporation method.  
The cDNA libraries constructed by this method are  
full-length enriched cDNA library."

BASE COUNT 85 a 48 c 59 g 60 t  
ORIGIN

Query Match 33.0%; Score 241.4; DB 14; Length 252;  
Best Local Similarity 99.6%; Pred. No. 1.5e-57;  
Matches 242; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 373 CTGGAATCTAGAGAAATCGAAGCCACCACATCATGAGAAATGTGGCGGCTGTTCAA 432  
|||||  
Db 1 CTGGAATCTAGAGAAATCGAAGCCACCACATCATGAGAAATGTGGCGGCTGTTCAA 60  
|||||

QY 433 AATGTCCCCCTGATTAAGGAGAAAGGCAACAGTAACTGACGTCCAGACATGAAGA 492  
|||||  
Db 61 AATGTCCCCCTGATTAAGGAGAAAGGCAACAGTAACTGACGTCCAGACATGAAGA 120  
|||||

QY 493 GATCGCAGTGCAGGAGCAAAATGCTTGTGTTGAAGAAAGTGAAGTCCCTG 552  
|||||

Db 121 GATGCCAGTCCACGAGCAAAATTCATTTGCTTTGTTAGAGAAAGTGAATCTTGG 180  
QY 553 CTCTTTGTGAGAAATTTTCATGAGATATGTTGGCCCAATAAAGATGACATTT 612  
|||||  
Db 181 CTCTTTGTGAGAAATTTTCATGAGATATGTTGGCCCAATAAAGATGACATTT 240  
|||||

QY 613 CAA 615  
|||  
Db 241 CAA 243

RESULT 12  
AA633404 188 bp mRNA linear EST 28-OCT-1997  
LOCUS np69h11.s1 NCI-CGAP Br2 Homo sapiens cDNA clone IMAGE:1131621 3'  
DEFINITION similar to SW:MIK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM  
CHANNEL PROTEIN ;, mRNA sequence.  
ACCESSION AA633404  
VERSION AA633404.1 GI:2555264  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 188)  
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
Unpublished (1997)  
JOURNAL Contact: Robert Strausberg, Ph.D.  
COMMENT Email: cgapbs-remail.nih.gov  
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.  
Emmert-Buck, M.D., Ph.D.  
CDNA Library Preparation: M. Bento Soares, Ph.D.  
DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
www.bio.llnl.gov/dbp/image/image.html  
Insert Length: 785 Std Error: 0.00  
Seq primer: -40m13 fwd. RT from Amersham  
High quality sequence stop: 167.  
Location/Qualifiers  
1..188  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1131621"  
/clone\_id="NCI-CGAP Br2"  
/sex="female, pooled"  
/issue\_type="breast"  
/lab\_host="DH10B"  
/note="Vector: pTV73D-Pac (Pharmacia) with a modified  
polylinker; Int strand cDNA was prepared from pooled bulk  
breast tumor tissue, and was then primed with a Not I -  
R1 adaptor (Pharmacia), digested with Not I and cloned  
into the Not I and Eco RI sites of the modified pTV73  
vector. This library is the normalized version of  
NCI-CGAP Br1.1. Library was constructed by Bento Soares  
and M. Fatima Bonaldo."

BASE COUNT 66 a 45 c 45 g 32 t  
ORIGIN

Query Match 25.7%; Score 188; DB 9; Length 188;  
Best Local Similarity 100.0%; Pred. No. 1.9e-42;  
Matches 188; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 286 TGTGAATTCACAGAGAGGAGACACTTCAATGACCCCTACACCACTAATTTAGAGCA 345  
|||||  
Db 1 TGTGAATTCACAGAGAGGAGACACTTCAATGACCCCTACACCACTAATTTAGAGCA 60  
|||||

QY 346 CTGCGAGGAATACAGAGGCAAAATCTGATCTAGAAATGCAAGGCCACCTTCA 405  
|||||

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Db      61 CTGGCAGGAAAGATCAAGAGCCAAATCTTGATCTAGAGAATCGAAGGCCACCACATCCA 120
Oy      406 TGAGAACATTTGGTGGCGGTGATGTCATGCCCCGATAGGAGAAAGGACGACG 465
Db      121 TGGAGAACATTTGGTGGCGGTGATGTCATGCCCCGATAGGAGAAAGGACGACG 180
Oy      466 CTACATC 473
Db      181 CTACATC 188

RESULT 13
LOCUS    BG938225
DEFINITION 1AB014L12 Bovine Abomasum cDNA library Bos taurus cDNA 5', mRNA
ACCESSION BG938225
VERSION   BG938225.1 GI:14337597
KEYWORDS EST.
SOURCE    EST.
ORGANISM  Bos taurus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
           Bovidae; Bovinae; Bos.
REFERENCE 1 (bases 1 to 351)
AUTHORS   Moore,S.S., Hansen,C., Li,C., Fu,A., Meng,Y. and Li,G.
TITLE      cDNA's from bovine abomasum tissue
JOURNAL    Unpublished (2001)
COMMENT    Contact: Dr. Stephen Moore
           . Beet Genomics Laboratory
           Dept of AFNS, University of Alberta
           410 Agri/For, Dept of AFNS, U of A, Edmonton, AB, T6G 2P5, Canada
           Tel: 780 492 0169
           Fax: 780 492 4265
           Email: smoores@afns.ualberta.ca
           The sequence best matches gb:AP001719 (Homo sapiens genomic DNA,
           chromosome 21q, section 63/105) in main database at high score of
           212.0 and E-value of 9e-53
PCR PRIMERS
FORWARD: M13 Forward
BACKWARD: M13 Reverse
Seq primer: T3 primer
High quality sequence stop: 351
POLY-A-NO.

FEATURES
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         1..351
            /location/Qualifiers
            /organism="Bos taurus"
            /db_xref="taxon:9913"
            /clone_lib="Bovine Abomasum cDNA library"
            /sex="Two males and one female mixed"
            /tissue_type="Gastrointestinal tissue (GIT)"
            /cell_type="Epithelial"
            /dev_stage="Young adult"
            /lab_host="XLI-BlueMR", strain"
            /note="Organ: Abomasum; Vector: Uni-ZAPXR; Site_1: EcoR
            I; Site_2: Xho I"

BASE COUNT  104 a      89 c      76 g      82 t
ORIGIN
Query Match      24.0%; Score 175.4; DB 13; Length 351;
Best Local Similarity 86.5%; Pred. No. 9.5e-39;
Matches 205; Conservative 0; Mismatches 31; Indels 1; Gaps 1;

```

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Db      235 CTGCAAGCAAGGTGATGCTGAGAACTTCTACTATGTCATCTGTACCTATGATG 294
Oy      242 ATGGAATGTTCTCTT CATCATCGTGGCCATCTCTGTAGCATGTGAATCCAA 237
Db      295 ATCGGAATGTTCTCTTCCATCATCTGTAGCATCTCTGTGTAGCAGCGTGAATCCAA 351

RESULT 14
LOCUS    AA935321
DEFINITION 121 bp mRNA linear EST 07-JUL-1998
ACCESSION AA935321
VERSION   AA935321.1 GI:3092478
KEYWORDS EST.
SOURCE    human.
ORGANISM  Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 121)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
           Unpublished (1997)
JOURNAL    Contact: Robert Strausberg, Ph.D.
           Email: cgaps-remail.nih.gov
           Tissue Procurement: Christopher A. Moskalko, M.D., Ph.D., Michael
           Emmert-Buck, M.D., Ph.D.
           cDNA Library Preparation: M. Bento Soares, Ph.D.
           cDNA Library Arrayed by: Greg Lennon, Ph.D.
           DNA Sequencing by: Washington University Genome Sequencing Center
           clone distribution: NCI-CGAP clone distribution information can be
           found through the I.M.A.G.E. Consortium/LIN at:
           www-bio.llnl.gov/bdrip/image/image.html

Trace considered overall poor quality
Insert length: 763 Std Error: 0.00
Seq primer: -40ml3 fwd. RT from Amersham
High quality sequence stop: 1.

FEATURES
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            /db_xref="taxon:9606"
            /clone_image="1571680"
            /clone_lib="NCI-CGAP_Gc4"
            /tissue_type="pooled germ cell tumors"
            /lab_host="DH10B"
            /note="Vector: pTZ19-D-Pac (Pharmacia) with a modified
            polylinker: 1st strand cDNA was prepared from 3 pooled
            germ cell tumors, and was then primed with a Not I -
            oligo(dT) primer. Double-stranded cDNA was ligated to Eco
            RI adaptors (Pharmacia), digested with Not I and cloned
            into the Not I and Eco RI sites of the modified pTZ19
            vector. Library is normalized. Library was constructed by
            Bento Soares and M. Fatima Bonaldo."

BASE COUNT  33 a      31 c      28 g      29 t
ORIGIN
Query Match      16.3%; Score 119.4; DB 9; Length 121;
Best Local Similarity 99.2%; Pred. No. 4.4e-23;
Matches 120; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 121 C 121

## RESULT 15

AM869303/c

LOCUS AM869303 311 bp mRNA linear EST 22-MAY-2000  
DEFINITION MR3-SN0067-240400-006-f11 SN0067 Homo sapiens cDNA, mRNA sequence.

AM869303

AM869303.1 GI:8003356

EST.

KEYWORDS

SOURCE

human.

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homindae; Homo.

1 (bases 1 to 311)

## REFERENCE

## AUTHORS

Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,  
Nagal,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,P.F.,  
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bala,G.S., Simpson,D.H.,  
Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare  
,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and  
Simpson,A.J.Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags

Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

## JOURNAL

## MEDLINE

## COMMENT

Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?cl=6t2-MR3-SN0067-240

400-006-f11&amp;t3=2000-04-24&amp;t4=1)

Seq primer: puc 18 forward

High quality sequence start: 11

High quality sequence stop: 74.

Location/Qualifiers

1..311

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone\_lib="SN0067"

/dev\_stage="Adult"

/note="Organ: stomach; normal; Vector: puc18; Site\_1: Sma1;  
Site\_2: Sma1; A mini-library was made by cloning products  
derived from ORESSES PCR (U.S. Letters Patent application  
No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the puc 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."

## BASE COUNT

60 a 80 c 76 g 95 t

## ORIGIN

Query Match 15.4%; Score 113; DB 10; Length 311;  
Best local Similarity 91.6%; Pred. No. 4.5e-21;

Matches 131; Conservative 0; Mismatches 10; Indels 2; Gaps 1;

QY 60 AGCAGAGGAGGAGCATGCTACTTATCCAAATTCACAGACGCTGGAGAGCTCTCC 119  
 |||  
 DB 196 AGCAGAGGAGGAGCATGCTACTTATCCAAATTCACAGACGCTGGAGAGCTCTCC 137  
 |||  
 QY 120 GAGGATTTTATTTACTTATATGACAATGG--CGCCAGACACAAACAGCTGAGCAAGA 177  
 |||  
 DB 136 GAGGATTTTATTTATCATTGACATCGGCGGCGAGACACGACAGCTGAGCGAGA 77  
 |||  
 QY 178 GGCCCTCCAGCCAAAGTTGATG 200  
 |||  
 DB 76 GGCCCTCCAGCCAAAGTTGATG 54

GenCore version 5.1.4.p5.4578  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - protein search, using sw model

Run on: May 15, 2003, 14:23:09 ; Search time 14 Seconds  
(without alignments)  
258.501 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632

Sequence: 1 MSTLSNFTQLEVDVFRRI...EESKATIHENIGAGFKMSP 123

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 262574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: /cgn2\_6/ptodata/1/1aa/5A.COMB.pep.\*  
2: /cgn2\_6/ptodata/1/1aa/5B.COMB.pep.\*  
3: /cgn2\_6/ptodata/1/1aa/6A.COMB.pep.\*  
4: /cgn2\_6/ptodata/1/1aa/6B.COMB.pep.\*  
5: /cgn2\_6/ptodata/1/1aa/PCTUS.COMB.pep.\*  
6: /cgn2\_6/ptodata/1/1aa/Backfile1.pep.\*

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	132.5	21.0	129	US-09-069-896-3	Sequence 3, Appli
2	132.5	21.0	129	US-09-135-021-78	Sequence 78, Appli
3	132.5	21.0	129	US-09-135-020-4	Sequence 4, Appli
4	132.5	21.0	129	US-09-135-010A-4	Sequence 4, Appli
5	132.5	21.0	129	US-09-444-871-4	Sequence 4, Appli
6	132.5	21.0	129	US-09-597-735-4	Sequence 4, Appli
7	132.5	21.0	129	US-09-444-295-4	Sequence 4, Appli
8	132.5	21.0	129	US-09-471-468-3	Sequence 4, Appli
9	132.5	21.0	129	US-09-597-732-4	Sequence 4, Appli
10	132.5	21.0	129	US-09-679-185-2	Sequence 3, Appli
11	132.5	21.0	132	US-08-118-101A-6	Sequence 6, Appli
12	128	20.3	130	US-09-069-896-4	Sequence 4, Appli
13	128	20.3	130	US-09-471-468-4	Sequence 4, Appli
14	127.5	20.2	129	US-09-679-185-4	Sequence 4, Appli
15	74	11.7	262	US-09-134-001C-5532	Sequence 5532, Ap
16	70	11.1	170	US-09-069-896-1	Sequence 1, Appli
17	70	11.1	170	US-09-471-468-1	Sequence 1, Appli
18	69	10.9	411	US-08-887-534A-80	Sequence 80, Appli
19	66.5	10.5	563	US-09-134-001C-3172	Sequence 3172, Ap
20	65.5	10.4	648	US-09-134-001C-5161	Sequence 5161, Ap
21	65.5	10.4	987	US-08-436-054-6	Sequence 6, Appli
22	65.5	10.4	987	US-08-436-054-6	Sequence 6, Appli
23	65.5	10.4	987	PCT-US95-08812-6	Sequence 6, Appli
24	65.5	10.4	1276	US-08-222-616-24	Sequence 24, Appli
25	65.5	10.4	1276	US-08-446-648-24	Sequence 24, Appli
26	65.5	10.4	1276	PCT-US95-04228-24	Sequence 24, Appli
27	65.5	10.4	2016	US-09-634-920-4	Sequence 4, Appli

28	65	10.3	1835	4	US-09-404-650-5	Sequence 5, Appli
29	65	10.3	2175	4	US-09-404-650-2	Sequence 2, Appli
30	65	10.3	2188	4	US-09-404-650-4	Sequence 4, Appli
31	64.5	10.2	970	2	US-08-673-789-7	Sequence 7, Appli
32	64.5	10.2	973	1	US-08-162-809-10	Sequence 10, Appli
33	64.5	10.2	976	4	US-09-302-812-4	Sequence 4, Appli
34	64.5	10.2	976	4	US-09-511-477-4	Sequence 4, Appli
35	64.5	10.2	976	4	US-09-511-507-4	Sequence 4, Appli
36	64	10.1	1367	2	US-08-249-687C-2	Sequence 2, Appli
37	64	10.1	1367	2	US-08-625-819-2	Sequence 2, Appli
38	64	10.1	1367	3	US-08-746-559A-2	Sequence 2, Appli
39	64	10.1	1367	4	US-08-864-641B-18	Sequence 18, Appli
40	63.5	10.0	300	4	US-09-293-549-14	Sequence 14, Appli
41	63.5	10.0	439	3	US-08-448-722A-5	Sequence 5, Appli
42	63.5	10.0	439	4	US-08-189-309B-5	Sequence 5, Appli
43	63.5	10.0	2332	1	US-08-276-894A-2	Sequence 2, Appli
44	63.5	10.0	2351	1	US-08-366-851A-2	Sequence 2, Appli
45	63.5	10.0	2351	6	5171844-2	Patent No. 5171844

#### ALIGNMENTS

RESULT 1  
US-09-069-896-3  
Sequence 3, Application US/09069896  
Patent No. 6071720  
GENERAL INFORMATION:  
APPLICANT: Hillman, Jennifer L.  
APPLICANT: Patterson, Chandra  
APPLICANT: Corley, Neil C.  
TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM  
TITLE OF INVENTION: CHANNEL HOMOLOG  
NUMBER OF SEQUENCES: 4  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Incyte Pharmaceuticals, Inc.  
STREET: 3174 Porter Drive  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94304  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FASTSD for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/069, 896  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Carrone, Michael C  
REGISTRATION NUMBER: 39,132  
REFERENCE/DOCKET NUMBER: PF-0507 US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-855-0555  
TELEFAX: 650-845-4166  
TELEX:  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 129 amino acids  
TYPE: amino acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
IMMEDIATE SOURCE:  
LIBRARY: Genbank  
CLONE: 452497  
US-09-069-896-3  
Query Match 21.0%; Score 132.5; DB 3; Length 129;





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; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-444-871-4

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Query Match	21.0%;	Score 132.5;	DB 4;	Length 129;
Best Local Similarity	45.1%;	Pred. No. 9.3e-08;		
Matches 23; Conservative	15;	Mismatches 12;	Indels 1;	Gaps 1;

OY 51 LYLMWIGMFSFLIIVLVSTVKSRRHSNDPHYQIYVED-WOEKYSOI 100  
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DB 45 LYLWMVLGFEGFETGLMLSYIRSKKLEHSNDPNNVLTESDAQWEKDRAV 95

RESULT 6  
US-09-597-735-4

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; Sequence 4, Application US/09597735
; Patent No. 6420124
; GENERAL INFORMATION:

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; APPLICANT: Keating, Mark T.
; APPLICANT: Sangulnnett1, Michael C.
; APPLICANT: Curran, Mark E.

```

; APPLICANT: Landes, Gregory M.  
; APPLICANT: Connors, Timothy D.  
; APPLICANT: Burn, Timothy C.

```

; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: KVL0T1 - A LONG QT SYNDROME GENE
; FILE REFERENCE: 2323-133
;

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;  
 ; CURRENT APPLICATION NUMBER: 05/09/591,,/355  
 ;  
 ; CURRENT FILING DATE: 2000-06-19  
 ;  
 ; EARLIER APPLICATION NUMBER: 09/135,010  
 ;  
 ; EARLIER FILING DATE: 1008-08-17

EARLIER FILING DATE: 1998-08-17  
EARLIER APPLICATION NUMBER: 60/094,477  
EARLIER FILING DATE: 1998-07-29  
EARLIER APPLICATION NUMBER: 08/931 068

EARLIER FILING DATE: 1997-08-29  
EARLIER APPLICATION NUMBER: 08/739,383  
EARLIER FILING DATE: 1996-10-29

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; EARLIER APPLICATION NUMBER: 60/019,014
; EARLIER FILING DATE: 1995-12-22
; NUMBER OF SEQ ID NOS: 116

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; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 129

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; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-597-735-4

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Query Match	21.0%	Score 132.5;	DB 4;	Length 129;
Best Local Similarity	45.1%;	Pred. No. 9.3e-08;		
Matches	23;	Conservative	12;	Totals
		Mismatches	1;	Calls
			1;	

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         ||:::| | : |:| ::|| :|||||: | | ||| | : :
Db      45 IYVI MWIGEEGETTGTMTSYTRSKRTFHSNDBENVVTESDAWQETRAVV 95
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## RESULT 7

US-09-444-295-4  
; Sequence 4, Application US/09444295  
; Patent No. 6432644

```

: GENERAL INFORMATION:
: APPLICANT: Keating, Mark T.
: APPLICANT: Sanquineti, Michael C

```

```

; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN mink WHICH
; TITLE OF INVENTION: CAUSE ARRHYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING

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; TITLE OF INVENTION: KCNE1 AS AN IQT GENE
; FILE REFERENCE: 2323-131
; CURRENT APPLICATION NUMBER: US/09/444,295

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; CURRENT FILING DATE: 1999-11-22

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1 PRIOR APPLICATION NUMBER: 09/135,020
2
3 PRIOR FILING DATE: 1998-08-17
4
5 PRIOR APPLICATION NUMBER: 08/921,068
6
7 PRIOR FILING DATE: 1997-08-29
8
9 PRIOR APPLICATION NUMBER: 08/733,383
10
11 PRIOR FILING DATE: 1996-10-29
12
13 PRIOR APPLICATION NUMBER: 60/019,014
14
15 PRIOR FILING DATE: 1995-12-22
16
17 PRIOR APPLICATION NUMBER: 60/094,477
18
19 PRIOR FILING DATE: 1998-07-29
20
21 NUMBER OF SEQ ID NOS: 114
22
23 SOFTWARE: PatentIn Ver. 2.0
24
25 SEQ ID NO: 4
26
27 LENGTH: 129
28
29 TYPE: prt
30
31 ORGANISM: Homo sapiens
32
33 IS-09-444-295-4

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Query Match	21.0%;	Score 132.5;	DB 4;	Length 129;
Best Local Similarity	45.1%;	Pred. No. 9.3e-08;		
Matches 23;	Conservative 15;	Mismatches 12;	Indels 1;	Gaps 1

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Oy 51 LYLMWIGMFSTIIVALLVSTVKSRRHESNDPHYIVED-WQEKYKSQI 1000
      |:::| | | :::| ::| |::|::| | | | | | | | : :
Db 45 LYVLWVGFEFGFTGLIMLSYIRSKLLEHSNDPNVYIESDAWQEKDKAY 95
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## RESULT 8

US-09-471-468-3  
; Sequence 3, Application US/09471468  
; Patent No. 6432687

```

; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Patterson, Chandra

```

APPLICANT: Corley, Neil C.  
TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM  
TITLE OF INVENTION: CHANNEL HOMOLOG  
NUMBER OF DRAWINGS: 4

NUMBER OF SEQUENCES: 4  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Incyte Pharmaceuticals, Inc.  
CITY: 3174 Darter Drive  
STATE: CA 94025  
COUNTRY: USA

STREET: 31/4  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA

COGNINI: USA  
ZIP: 94304  
COMPUTER READABLE FORM:  
MEDITIM TYPE: D4skott

; MEDIUM LIFE: DISKETTE  
 ; COMPUTER: IBM Compatible  
 ; OPERATING SYSTEM: DOS  
 ; SOFTWARE: FASTSEQ for W

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CURRENT APPLICATION DATA:
APPLICATION NUMBER:  US/09/471,468
FILING DATE:

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; FILING DATE:
;
; CLASSIFICATION:
;
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/069,896
;

```

FILED DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Cerrone, Michael C

REGISTRATION NUMBER: 39,132  
REFERENCE/DOCKET NUMBER: PF-0507 US  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-855-0555  
TELEFAX: 650-845-4166  
TELEX:

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; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 129 amino acids

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; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
;

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; IMMEDIATE SOURCE:
; LIBRARY: GenBank

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: CLONE: 452497
US-09-471-468-3

Query Match      21.0%; Score 132.5; DB 4; Length 129;
Best Local Similarity 45.1%; Pred. No. 9.3e-08;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLWMIQMFSEIIVAILVSTVKSRRRHSNDPRHYQIYVED-WQEKYSQI 100
||||:| | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 45 LYLWVLGFGFETLIGIMLSYIRSKLEHSNDPENVYIESDAMQEKRAVY 95

RESULT 9
US-09-597-732-4
: Sequence 4, Application US/09597732
: Patent No. 6451534
: GENERAL INFORMATION:
: APPLICANT: Keating, Mark T.
: APPLICANT: Sangunetti, Michael C.
: APPLICANT: Curran, Mark E.
: APPLICANT: Landes, Gregory M.
: APPLICANT: Connors, Timothy D.
: APPLICANT: Burn, Timothy C.
: APPLICANT: Splawski, Igor
: TITLE OF INVENTION: KVLQTL - A LONG QT SYNDROME GENE
: FILE REFERENCE: 2323-133
: CURRENT APPLICATION NUMBER: US/09/597,732
: CURRENT FILING DATE: 2000-06-19
: PRIOR APPLICATION NUMBER: 09/135,010
: PRIOR FILING DATE: 1998-08-17
: PRIOR APPLICATION NUMBER: 60/094,477
: PRIOR FILING DATE: 1998-07-29
: PRIOR APPLICATION NUMBER: 08/921,068
: PRIOR FILING DATE: 1997-08-29
: PRIOR APPLICATION NUMBER: 08/739,383
: PRIOR FILING DATE: 1996-10-29
: PRIOR APPLICATION NUMBER: 60/019,014
: PRIOR FILING DATE: 1995-12-22
: NUMBER OF SEQ ID NOS: 116
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 4
: LENGTH: 129
: TYPE: PRT
: ORGANISM: Homo sapiens
US-09-597-732-4

Query Match      21.0%; Score 132.5; DB 4; Length 129;
Best Local Similarity 45.1%; Pred. No. 9.3e-08;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYLWMIQMFSEIIVAILVSTVKSRRRHSNDPRHYQIYVED-WQEKYSQI 100
||||:| | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 45 LYLWVLGFGFETLIGIMLSYIRSKLEHSNDPENVYIESDAMQEKRAVY 95

RESULT 10
US-09-679-185-2
: Sequence 2, Application US/09679185
: Patent No. 6458542
: GENERAL INFORMATION:
: APPLICANT: George Jr., Alfred L.
: APPLICANT: Roden, Dan M
: TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO
: FILE REFERENCE: ATTORNEY DOCKET NO. 6458542 1242-33-2
: CURRENT APPLICATION NUMBER: US/09/679,185
: CURRENT FILING DATE: 2000-10-04
: PRIOR APPLICATION NUMBER: 60/158,696
: PRIOR FILING DATE: 1999-10-08
: NUMBER OF SEQ ID NOS: 11
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 2
: LENGTH: 129

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: TYPE: PRT
: ORGANISM: Homo sapiens
US-09-679-185-2

Query Match                               21.0%; Score 132.5; DB 4; Length 129;
Best Local Similarity 45.1%; Pred. No. 9.3e-08;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYTMVMIGMSFTIVALLVTVKSRKRRHSNDPYHQYIVED-WQEKYSQI 100
||||:| | | : ||:| | | | | | | | | | | | | | | | | | | | |
DB 45 LYTMVIGFGFPTLGLIMLSYIRSKLRLHSNDPENVYIESDAMQEKDAY 95

RESULT 11
US-08-118-101A-6
: Sequence 6, Application US/08118101A
: Patent No. 5620892
: GENERAL INFORMATION:
: APPLICANT: Kurtz, Stephen E.
: APPLICANT: Knickerbocker, Aron M.
: APPLICANT: McCullough, John R.
: TITLE OF INVENTION: A STRAIN OF SACCHAROMYCES CEREVISIAE
: NUMBER OF SEQUENCES: 16
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Burton Rodney
: STREET: P.O. Box 4000
: CITY: Princeton
: STATE: New Jersey
: COUNTRY: U.S.A.
: ZIP: 08543-4000
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/118,101A
: FILING DATE:
: CLASSIFICATION: 435
: ATTORNEY/AGENT INFORMATION:
: NAME: Gaul, Timothy J.
: REGISTRATION NUMBER: 33,111
: REFERENCE/DOCKET NUMBER: DC/7
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (609) 252-5901
: TELEFAX: (609) 252-4526
: INFORMATION FOR SEQ ID NO: 6:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 132 amino acids
: TYPE: amino acid
: TOPOLOGY: linear
: MOLECULE TYPE: protein
US-08-118-101A-6

Query Match                               21.0%; Score 132.5; DB 1; Length 132;
Best Local Similarity 45.1%; Pred. No. 9.6e-08;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

OY 51 LYTMVMIGMSFTIVALLVTVKSRKRRHSNDPYHQYIVED-WQEKYSQI 100
||||:| | | : ||:| | | | | | | | | | | | | | | | | | | | |
DB 48 LYTMVIGFGFPTLGLIMLSYIRSKLRLHSNDPENVYIESDAMQEKDAY 98

RESULT 12
US-09-069-896-4
: Sequence 4, Application US/09069896
: Patent No. 6071720
: GENERAL INFORMATION:
: APPLICANT: Hillman, Jennifer L.
: APPLICANT: Patterson, Chandra
: APPLICANT: Corley, Neil C.
: TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM

```

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; TITLE OF INVENTION: CHANNEL HOMOLOG
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
;
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/069,896
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Cerrone, Michael C
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0507 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; TELEX:
;
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 130 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 203977
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; US-09-069-896-4
;
; Query Match 20.3%; Score 128; DB 3; Length 130;
; Best Local Similarity 41.4%; Pred. No. 3e-07;
; Matches 29; Conservative 17; Mismatches 20; Indels 4; Gaps 3;
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; QY 51 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKS--QILNLEESK 107
; ||:::| | | | |:::| | | | |:::| | | | |:::| | | | |:::| | | | |
; DB 46 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKS--QILNLEESK 105
;
; QY 108 AT-IHENIGA 116
; | : | | |
; DB 106 ACYVIENQAA 115
;
; RESULT 13
; US-09-471-468-4
; Sequence 4, Application US/09471468
; Patent No. 6432687
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Patterson, Chandra
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM
; TITLE OF INVENTION: CHANNEL HOMOLOG
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
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; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/471,468
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/069,896
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Cerrone, Michael C
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0507 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-855-0555
; TELEFAX: 650-845-4166
; TELEX:
;
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 130 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 203977
;
; US-09-471-468-4
;
; Query Match 20.3%; Score 128; DB 4; Length 130;
; Best Local Similarity 41.4%; Pred. No. 3e-07;
; Matches 29; Conservative 17; Mismatches 20; Indels 4; Gaps 3;
;
; QY 51 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKS--QILNLEESK 107
; ||:::| | | | |:::| | | | |:::| | | | |:::| | | | |:::| | | | |
; DB 46 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKS--QILNLEESK 105
;
; QY 108 AT-IHENIGA 116
; | : | | |
; DB 106 ACYVIENQAA 115
;
; RESULT 14
; US-09-679-185-4
; Sequence 4, Application US/09679185
; Patent No. 6458542
; GENERAL INFORMATION:
; APPLICANT: Roden, Dan M
; APPLICANT: George Jr., Alfred L.
; TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO
; TITLE OF INVENTION: DRUG-INDUCED CARDIAC ARYTHMIA
; FILE REFERENCE: Attorney Docket No. 6458542 1242-33-2
; CURRENT APPLICATION NUMBER: US/09/679,185
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: 60/158,696
; PRIOR FILING DATE: 1999-10-08
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 4
; LENGTH: 129
; TYPE: PRT
; ORGANISM: Homo sapiens
;
; US-09-679-185-4
;
; Query Match 20.2%; Score 127.5; DB 4; Length 129;
; Best Local Similarity 43.1%; Pred. No. 3.4e-07;
; Matches 22; Conservative 16; Mismatches 12; Indels 1; Gaps 1;
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; QY 51 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKSQI 100
; ||:::| | | | |:::| | | | |:::| | | | |:::| | | | |:::| | | | |
; DB 45 LYLAMWIGMFSPIVALIVSTVSKRREHSDNPYHQIYVED-WQEKYKSQI 95
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Result No.	Score	Query Match	Length	DB	ID	Description
1	730.4	99.8	732	9	US-10-000-151B-5	Sequence 5, Appl1
2	730.4	99.8	113604	9	US-10-227-195A-1	Sequence 1, Appl1
3	730.4	99.8	113604	9	US-10-227-195A-2	Sequence 2, Appl1
4	372.5	50.8	372	10	US-09-864-761-33139	Sequence 3, Appl1
5	368	50.3	3	10	US-09-864-761-3463	Sequence 3663, Appl1
6	312	42.6	312	10	US-09-864-761-20233	Sequence 20233, Appl1
7	306	41.8	471	10	US-09-864-761-16671	Sequence 16671, Appl1
8	53.2	7.3	231	10	US-09-864-761-20783	Sequence 20783, Appl1
9	53.2	7.3	380	10	US-09-864-761-17593	Sequence 17593, Appl1
10	53.2	7.3	1703	9	US-10-138-316-3	Sequence 3, Appl1
11	53.2	7.3	11604	9	US-10-227-195A-1	Sequence 1, Appl1
12	53.2	7.3	113604	9	US-10-227-195A-2	Sequence 2, Appl1
13	50.8	6.9	468	10	US-09-864-761-4026	Sequence 4026, Appl1
14	46	6.3	381	10	US-09-864-761-810	Sequence 810, Appl1
15	36	4.9	1146	10	US-09-853-386-111	Sequence 111, Appl1
16	36	4.9	1206	10	US-09-853-386-107	Sequence 107, Appl1
17	36	4.9	1206	10	US-09-853-386-117	Sequence 117, Appl1
18	36	4.9	1485	10	US-09-191-724-1	Sequence 1, Appl1
19	36	4.9	1562	10	US-09-191-724-14	Sequence 14, Appl1

5	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
6	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
7	33.6	4.6	653	9	US-10-184-654-402	Sequence 402, App
8	33.6	4.6	653	9	US-10-184-654-402	Sequence 402, App
9	33.6	4.6	1140	12	US-10-003-356-9	Sequence 6, App1
10	33.6	4.6	2781	12	US-10-003-356-9	Sequence 9, App1
11	33.4	4.6	4104	9	US-09-992-598-277	Sequence 277, App
12	33.4	4.6	4104	9	US-09-988-283A-277	Sequence 277, App
13	33.4	4.6	4104	9	US-09-988-735-277	Sequence 277, App
14	33.4	4.6	4104	9	US-09-990-444-277	Sequence 277, App
15	33.4	4.6	4104	9	US-09-989-730-277	Sequence 277, App
16	33.4	4.6	4104	9	US-09-990-436-277	Sequence 277, App
17	33.4	4.6	4104	9	US-09-991-181-277	Sequence 277, App
18	33.4	4.6	4104	9	US-09-993-687-277	Sequence 277, App
19	33.4	4.6	4104	9	US-09-988-734-277	Sequence 277, App
20	33.4	4.6	4104	9	US-10-028-072-449	Sequence 449, App
21	33.4	4.6	4104	9	US-09-997-653-277	Sequence 277, App
22	33.4	4.6	4104	9	US-09-993-667-277	Sequence 277, App
23	33.4	4.6	4104	9	US-10-121-049-449	Sequence 449, App
24	33.4	4.6	4104	9	US-10-123-904-449	Sequence 449, App
25	33.4	4.6	4104	9	US-10-140-470-449	Sequence 449, App
26	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
27	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
28	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
29	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
30	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
31	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
32	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
33	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
34	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
35	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
36	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
37	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
38	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
39	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
40	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
41	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
42	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
43	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
44	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App
45	33.4	4.6	4104	9	US-09-990-438-277	Sequence 277, App

## ALIGNMENTS

```

RESULT 1
US-10-000-151b-5
: Sequence 5, Application US/10000151B
: Publication No. US20030013136A1
: GENERAL INFORMATION:
: APPLICANT: Balser, Jeffrey R.
: APPLICANT: George, Alfred L.
: TITLE OF INVENTION: HUMAN KRII REGULATION OF HERP POTASSIUM CHANNEL BLOCK
: FILE REFERENCE: Vandebuilt Ref No. US20030013136A1 VU0120; Attorney Docket No. US-
: CURRENT APPLICATION NUMBER: US/10/000.151B
: CURRENT FILING DATE: 2000-10-30
: NUMBER OF SEQ ID NOS: 5
: SOFTWARE: PatentIn version 3.1
: SEQ ID NO 5
: LENGTH: 732
: TYPE: DNA
: ORGANISM: Homo sapiens
: US-10-000-151b-5

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Query Match	99.8%	Score 730.4	DB 9	Length 732
Best Local Similarity	99.9%	Pred. No. 4,6e-215		
Matches 731	Conservative	0	Mismatches 1	Indels 0
			Gaps	0
QY	1	CAATCCAGAAAAGATCCGTTTCCCTAACCTGTTCCGCTATTTTATTTATTTAAATGCA	60	
Db	1	CAATCCAGAAAAGATCCGTTTCCCTAACCTGTTCCGCTATTTTATTTATTTAAATGCA	60	
QY	61	GCAGAGAGGAGAGCATCTTACTTTATTCATAATTCCACAGACGGTGGAGAAGCGTCCG	120	
Db	61	GCAGAGAGGAGAGCATCTTACTTTATTCATAATTCCACAGACGGTGGAGAAGCGTCCG	120	
QY	121	AAGGATTTTTTAACTTATATGACAAATTGGGCCGACACAAACAGCTGAGCAAGGC	180	
Db	121	AAGGATTTTTTAACTTATATGACAAATTGGGCCGACACAAACAGCTGAGCAAGGC	180	
QY	181	CCGCCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGSGTAT	240	
Db	181	CCGCCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGSGTAT	240	
QY	241	GATTGGAAAGTCTCTTTTCATCATGCTGGCCATCTGCTGAGCACTGTGAATCCAAAG	300	
Db	241	GATTGGAAAGTCTCTTTTCATCATGCTGGCCATCTGCTGAGCACTGTGAATCCAAAG	300	

QY 301 ACGGAAACATCCATGATGACCCCTACACAGTACATTTGTAGAGAGCTGGCAGAGAAAGTA 360  
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Db 301 ACGGAAACATCCATGATGACCCCTACACAGTACATTTGTAGAGAGCTGGCAGAGAAAGTA 360  
QY 361 CAAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGC 420  
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Db 361 CAAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGC 420  
QY 421 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGGACCAAGTACATCTGAGCTC 480  
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Db 421 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGGACCAAGTACATCTGAGCTC 480  
QY 481 CAGACATGAAAGAGATGCGCAGTGCACGAGGCAATCCAAATGTCTTGTAGAGAAA 540  
|||||  
Db 481 CAGACATGAAAGAGATGCGCAGTGCACGAGGCAATCCAAATGTCTTGTAGAGAAA 540  
QY 541 GTGAGTTCCTGCTCTCTTTGTAGAAATTTTCATGAGATTAATGTGGTGGCCAAATTAAGA 600  
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Db 541 GTGAGTTCCTGCTCTCTTTGTAGAAATTTTCATGAGATTAATGTGGTGGCCAAATTAAGA 600  
QY 601 TAGATGACATTTCAATCTCAAGTATTTATGCTTCTGTTGAGCAATATTTTGTGCTGA 660  
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Db 601 TAGATGACATTTCAATCTCAAGTATTTATGCTTCTGTTGAGCAATATTTTGTGCTGA 660  
QY 661 AGACCTCTTTACTTTCGGGCAAGTGAATGTCATTTTATCAATATCATGATGAAAT 720  
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Db 661 AGACCTCTTTACTTTCGGGCAAGTGAATGTCATTTTATCAATATCATGATGAAAT 720  
QY 721 AAAGCCAAATTT 732  
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Db 721 AAAGCCAAATTT 732

RESULT 2  
US-10-227-195A-1  
; Sequence 1, Application US/10227195A  
; Publication No. US2003007633A1  
; GENERAL INFORMATION:  
; APPLICANT: Cox, David  
; APPLICANT: Arnold, Deana  
; TITLE OF INVENTION: Haplotype structure of chromosome 21  
; FILE REFERENCE: 1030U1  
; CURRENT APPLICATION NUMBER: US/10/227,195A  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1  
; LENGTH: 113604  
; TYPE: DNA  
; ORGANISM: Human  
; NAME/KEY: misc.feature  
; LOCATION: 7175, 7204, 36973, 66372, 76921, 81512, 88727  
; OTHER INFORMATION: n = G or C  
US-10-227-195A-1

Query Match 99.8%; Score 730.4; DB 9; Length 113604;  
Best Local Similarity 99.9%; Pred. No. 1.5e-213;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAATCCAGAAAAGATCCGTTTCTTAACCTGTGCGCTATTTTATTAATTTAAATTGCA 60  
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Db 17403 CAATCCAGAAAAGATCCGTTTCTTAACCTGTGCGCTATTTTATTAATTTAAATTGCA 17462  
QY 61 GCAGAGGAGAGCATGCTCTTATTCATTCACACAGCGTGAAGACGCTCCG 120  
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Db 17463 GCAGAGGAGAGCATGCTCTTATTCATTCACACAGCGTGAAGACGCTCCG 17522  
QY 121 AAGGATTTTATTTACTTATATGAGCAATTTGGCGCAGAACACAAACAGCTGAGCAAGAGG 180  
|||||  
Db 17523 AAGGATTTTATTTACTTATATGAGCAATTTGGCGCAGAACACAAACAGCTGAGCAAGAGG 17582

QY 181 CTTCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTTACCTAGTGTAT 240  
|||||  
Db 17583 CTTCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTTACCTAGTGTAT 17642  
QY 241 GATTGGAATTTCTTTTATCATCTGTTGGCATCTCTGTGAGCATCTGTGAATCCAGAG 300  
|||||  
Db 17643 GATTGGAATTTCTTTTATCATCTGTTGGCATCTCTGTGAGCATCTGTGAATCCAGAG 17702  
QY 301 ACGGAAACATCCATGATGACCCCTACACAGTACATTTGTAGAGAGCTGGCAGAGAAAGTA 360  
|||||  
Db 17703 ACGGAAACATCCATGATGACCCCTACACAGTACATTTGTAGAGAGCTGGCAGAGAAAGTA 17762  
QY 361 CAAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGC 420  
|||||  
Db 17763 CAAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGC 17822  
QY 421 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGGACCAAGTACATCTGAGCTC 480  
|||||  
Db 17823 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGGACCAAGTACATCTGAGCTC 17882  
QY 481 CAGACATGAAAGAGATGCGCAGTGCACGAGGCAATCCAAATGTCTTGTAGAGAAA 540  
|||||  
Db 17883 CAGACATGAAAGAGATGCGCAGTGCACGAGGCAATCCAAATGTCTTGTAGAGAAA 17942  
QY 541 GTGAGTTCCTGCTCTCTTTGTAGAAATTTTCATGAGATTAATGTGGTGGCCAAATTAAGA 600  
|||||  
Db 17943 GTGAGTTCCTGCTCTCTTTGTAGAAATTTTCATGAGATTAATGTGGTGGCCAAATTAAGA 18002  
QY 601 TAGATGACATTTCAATCTCAAGTATTTATGCTTCTGTTGAGCAATATTTTGTGCTGA 660  
|||||  
Db 18003 TAGATGACATTTCAATCTCAAGTATTTATGCTTCTGTTGAGCAATATTTTGTGCTGA 18062  
QY 661 AGACCTCTTTACTTTCGGGCAAGTGAATGTCATTTTATCAATATCATGATGAAAT 720  
|||||  
Db 18063 AGACCTCTTTACTTTCGGGCAAGTGAATGTCATTTTATCAATATCATGATGAAAT 18122  
QY 721 AAAGCCAAATTT 732  
|||||  
Db 18123 AAAGCCAAATTT 18134

RESULT 3  
US-10-227-195A-2  
; Sequence 2, Application US/10227195A  
; Publication No. US2003007633A1  
; GENERAL INFORMATION:  
; APPLICANT: Cox, David  
; APPLICANT: Arnold, Deana  
; TITLE OF INVENTION: Haplotype structure of chromosome 21  
; FILE REFERENCE: 1030U1  
; CURRENT APPLICATION NUMBER: US/10/227,195A  
; NUMBER OF SEQ ID NOS: 2  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 2  
; LENGTH: 113604  
; TYPE: DNA  
; ORGANISM: Human  
US-10-227-195A-2

Query Match 99.8%; Score 730.4; DB 9; Length 113604;  
Best Local Similarity 99.9%; Pred. No. 1.5e-213;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAATCCAGAAAAGATCCGTTTCTTAACCTGTGCGCTATTTTATTAATTTAAATTGCA 60  
|||||  
Db 17403 CAATCCAGAAAAGATCCGTTTCTTAACCTGTGCGCTATTTTATTAATTTAAATTGCA 17462  
QY 61 GCAGAGGAGAGCATGCTCTTATTCATTCACACAGCGTGAAGACGCTCCG 120  
|||||  
Db 17463 GCAGAGGAGAGCATGCTCTTATTCATTCACACAGCGTGAAGACGCTCCG 17522  
QY 121 AAGGATTTTATTTACTTATATGAGCAATTTGGCGCAGAACACAAACAGCTGAGCAAGAGG 180

Db	17523	AAGATTTTTATTACTAATATATGAGCAATTGGGCGCCAGAAACACAAGAGTGGACCAAGAGGC	17582
Qy	181	CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCAATCTCTGTACTCTATGCTGAT	240
Db	17583	CCCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCAATCTCTGTACTCTATGCTGAT	17642
Qy	241	GATTGGAATGTCTCTTTCATCATCTGGCCATCCTGGGAGACACTGTAATTCACAAG	300
Db	17643	GATTGGAATGTCTCTTTCATCATCTGGCCATCCTGGGAGACACTGTAATTCACAAG	17702
Qy	301	ACGGGAACACTCCAAATGACCCCTACCAACAGTACATTTGAGAGGACTGGCAGAAAAAGTA	360
Db	17703	ACGGGAACACTCCAAATGACCCCTACCAACAGTACATTTGAGAGGACTGGCAGAAAAAGTA	17762
Qy	361	CAAGAGCCAAATCTGTAATATAGAAATCGAAGGCGCACATCCATGAGAAATATGGTGC	420
Db	17763	CAAGAGCCAAATCTGTAATATAGAAATCGAAGGCGCACATCCATGAGAAATATGGTGC	17822
Qy	421	GGCTGGTTCCAAATGTGTCCTGATTAAGGAGAAAGGACCAAGTAACTACATGACGTC	480
Db	17823	GGCTGGTTCCAAATGTGTCCTGATTAAGGAGAAAGGACCAAGTAACTACATGACGTC	17882
Qy	481	CAGACATGAAGAGATGCGCATGCGCAGAGGCAATCCAAATGTCTTCTTCTTGAAGAAA	540
Db	17883	CAGACATGAAGAGATGCGCATGCGCAGAGGCAATCCAAATGTCTTCTTCTTGAAGAAA	17942
Qy	541	GTGAGTTCCTGCTCTTGTGAGAAATTTTCATGAGATTAATGTGGTGGCCCAATAGA	600
Db	17943	GTGAGTTCCTGCTCTTGTGAGAAATTTTCATGAGATTAATGTGGTGGCCCAATAGA	18002
Qy	601	TGATGACATTTCAATCTCAGTGAATTTATGCTTGTGCTTGGAGCAATATTTTGCTGA	660
Db	18003	TGATGACATTTCAATCTCAGTGAATTTATGCTTGTGCTTGGAGCAATATTTTGCTGA	18062
Qy	661	AGACCTCTTTACTCTTCCGGGCAAGTAAATGATTTTATCAATCAATGATGAAAAAT	720
Db	18063	AGACCTCTTTACTCTTCCGGGCAAGTAAATGATTTTATCAATCAATGATGAAAAAT	18122
Qy	721	AAAGCCAAATTT 732	
Db	18123	AAAGCCAAATTT 18134	

[illegible]

```

1 GENERAL INFORMATION:
2 APPLICANT: Penn, Sharon G.
3 APPLICANT: Rank, David R.
4 APPLICANT: Hanzel, David K.
5 APPLICANT: Chen, Wensheng
6 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
7 TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
8 FILE REFERENCE: Aeomica-X-1
9 CURRENT APPLICATION NUMBER: US 09/864,761
10 CURRENT FILING DATE: 2001-05-23
11 PRIOR APPLICATION NUMBER: US 60/180,312
12 PRIOR FILING DATE: 2000-02-04
13 PRIOR APPLICATION NUMBER: US 60/207,456
14 PRIOR FILING DATE: 2000-05-26
15 PRIOR APPLICATION NUMBER: US 09/632,366
16 PRIOR FILING DATE: 2000-08-03
17 PRIOR APPLICATION NUMBER: GB 24263.6
18 PRIOR FILING DATE: 2000-10-04
19 PRIOR APPLICATION NUMBER: US 60/236,359
20 PRIOR FILING DATE: 2000-09-27
21 PRIOR APPLICATION NUMBER: PCT/US01/00666
22 PRIOR FILING DATE: 2001-01-30
23 PRIOR APPLICATION NUMBER: PCT/US01/00667
24 PRIOR FILING DATE: 2001-01-30
25 PRIOR APPLICATION NUMBER: PCT/US01/00664
26 PRIOR FILING DATE: 2001-01-30
27 PRIOR APPLICATION NUMBER: PCT/US01/00669
28 PRIOR FILING DATE: 2001-01-30
29 PRIOR APPLICATION NUMBER: PCT/US01/00665
30 PRIOR FILING DATE: 2001-01-30
31 PRIOR APPLICATION NUMBER: PCT/US01/00668
32 PRIOR FILING DATE: 2001-01-30
33 PRIOR APPLICATION NUMBER: PCT/US01/00663
34 PRIOR FILING DATE: 2001-01-30
35 PRIOR APPLICATION NUMBER: PCT/US01/00662
36 PRIOR FILING DATE: 2001-01-30
37 PRIOR APPLICATION NUMBER: PCT/US01/00661
38 PRIOR FILING DATE: 2001-01-30
39 PRIOR APPLICATION NUMBER: PCT/US01/00670
40 PRIOR FILING DATE: 2001-01-30
41 PRIOR APPLICATION NUMBER: US 60/234,687
42 PRIOR FILING DATE: 2000-09-21
43 PRIOR APPLICATION NUMBER: US 09/608,408
44 PRIOR FILING DATE: 2000-06-30
45 PRIOR APPLICATION NUMBER: US 09/774,203
46 PRIOR FILING DATE: 2001-01-29
47 NUMBER OF SEQ ID NOS: 49117
48 SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
49 SEQ ID NO 3463
50 LENGTH: 450
51 TYPE: DNA
52 ORGANISM: Homo sapiens
53 FEATURE:
54 OTHER INFORMATION: MAP TO AP000052.1
55 OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
56 OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.1
57 OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3
58 OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.92
59 OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.4
60 OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
61 OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
62 OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.2
63 OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
64 OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
65 US-09-864-761-3463

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Query Match	Score	DB ID	Length
50.38	368	10	450
100.00	368	10	450

Matches 368; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CAATCCAGAAAGATCCGTTTTCTTAACCTGTGCGCTATTTATTTAAATTGCA 60  
|||||  
Db 83 CAATCCAGAAAGATCCGTTTTCTTAACCTGTGCGCTATTTATTTAAATTGCA 142

QY	61	GCAGAGGGAAGCAATGTCTACTTCTTATCCATTTCCACACAGACGGCTGGAAGACGTCTTCG	120
Db	143	GCAGAGGGAAGCAATGTCTACTTCTTATCCATTTCCACACAGACGGCTGGAAGACGTCTTCG	202
QY	121	AAGGATTTTATATCTTATATATGACAAATTGGCGCCAGAACCAACAGCTGAGCAAGGCC	180
Db	203	AAGGATTTTATATCTTATATATGACAAATTGGCGCCAGAACCAACAGCTGAGCAAGGCC	262
QY	181	CCTCCAGCCAAAGTGTATGCTGAGAACTTCTACTATGTCTCTGTACTCTTATCTGAT	240
Db	263	CCTCCAGCCAAAGTGTATGCTGAGAACTTCTACTATGTCTCTGTACTCTTATCTGAT	322
QY	241	GATTGGAATGTCTCTTCTCATCATCTGCGCATCTCTGGTGAAGCACTGAAATCCAAAG	300
Db	323	GATTGGAATGTCTCTTCTCATCATCTGCGCATCTCTGGTGAAGCACTGAAATCCAAAG	382
QY	301	ACGGGAACACTCCATGATGACCCCTTACACACAGTACATGTGTAGAGGACTGGCAGAAAAGTA	360
Db	383	ACGGGAACACTCCATGATGACCCCTTACACACAGTACATGTGTAGAGGACTGGCAGAAAAGTA	442
QY	361	CAAGAGCC 368	
Db	443	CAAGAGCC 450	



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; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 20233
; LENGTH: 312
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000052.1
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.92
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.2
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
; OTHER INFORMATION: EST_HUMAN HIT: A1246239.1, EVALUO 0.00e+00
; OTHER INFORMATION: SWISSPROT HIT: Q9Y6J6, EVALUO 3.00e-55
; OTHER INFORMATION: NT HIT: AF302095.1, EVALUO 0.00e+00
; US-09-864-761-20233

Query Match          42.6%; Score 312; DB 10; Length 312;
Best Local Similarity 100.0%; Pred. No. 5e-86;
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 127 TTTTATTACTTATATGCAATTTGGCCGAGAACACACAGCTGAGCAGAGCCCTCCA 186
      |||||||
DB 1 TTTTATTACTTATATGCAATTTGGCCGAGAACACACAGCTGAGCAGAGCCCTCCA 60

QY 187 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGTAACCTCATGATGATTTGG 246
      |||||||
DB 61 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGTAACCTCATGATGATTTGG 120

QY 247 AATGTCCTCTTCATCATGCTGGCCATCCTCTGCTGAGCACTGTGGAATCCAAAGACGGGA 306
      |||||||
DB 121 AATGTCCTCTTCATCATGCTGGCCATCCTCTGCTGAGCACTGTGGAATCCAAAGACGGGA 180

QY 307 ACACCTCAATGACCCCTACACACAGTACTTGTAGAGACTGGCAGAAAGTACAAAG 366
      |||||||
DB 181 ACACCTCAATGACCCCTACACACAGTACTTGTAGAGACTGGCAGAAAGTACAAAG 240

QY 367 CCAATCTTGAATCTTGAAGAATCGAAGGCCACCATCATGAGAACTTTGGCGGCTGG 426
      |||||||
DB 241 CCAATCTTGAATCTTGAAGAATCGAAGGCCACCATCATGAGAACTTTGGCGGCTGG 300

QY 427 GTTCAAAATGTC 438
      |||||||
DB 301 GTTCAAAATGTC 312

RESULT 7
US-09-864-761-16671
; Sequence 16671, Application US/09864761
; Patent No. US2002048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME- DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aemica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
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; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 16671
; LENGTH: 471
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000120.1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.98
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 0.67
; US-09-864-761-16671

Query Match          41.8%; Score 306; DB 10; Length 471;
Best Local Similarity 100.0%; Pred. No. 4.6e-84;
Matches 306; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAAATCCAGAAAAGATCCGTTTCTCTACCTGTTCGCTATTTTATTTAATTAATGCA 60
      |||||||
DB 166 CAAATCCAGAAAAGATCCGTTTCTCTACCTGTTCGCTATTTTATTTAATTAATGCA 225

QY 61 GCAGAGGGAAGCATGTCTACTTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCG 120
      |||||||
DB 226 GCAGAGGGAAGCATGTCTACTTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCG 285

QY 121 AAGGATTTTATTTACTTATATGACAAATTTGGCCGACACACAGAGCTGAGCAAGAGC 180
      |||||||
DB 286 AAGGATTTTATTTACTTATATGACAAATTTGGCCGACACACAGAGCTGAGCAAGAGC 345

QY 181 CCTCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATTCCTGATCCATCAAGTGAT 240
      |||||||
DB 346 CCTCAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATTCCTGATCCATCAAGTGAT 405

QY 241 GATTGGAATGTTCTTTTCATCATGCTGGCCATCCTGGTGAACACTGTGAATCCAGAG 300
      |||||||
DB 406 GATTGGAATGTTCTTTTCATCATGCTGGCCATCCTGGTGAACACTGTGAATCCAGAG 465

QY 301 ACGGGA 306
      |||||||
DB 466 ACGGGA 471
```

```

RESULT 8
US-09-864-761-20783/C
Sequence 20783, Application US/09864761
Patent No. US20020046763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
FILE REFERENCE: Aecm1ca-X-1
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US/09/864,761
PRIOR FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263,6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
SEQ ID NO 20783
LENGTH: 231
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000121.1
OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.1
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 0.94
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.59
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.74
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.66
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 0.66
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.7
OTHER INFORMATION: SWISSPROT HIT: P15382, EVALUATE 2.00e-39
OTHER INFORMATION: EST_HUMAN HIT: AW847275.1, EVALUATE 5.60e-01
US-09-864-761-20783
Query Match 7.3%, Score 53.2, DB 10, Length 231;
Best Local Similarity 63.6%, Pred No. 3e-06;
Matches 98, Conservative 0, Mismatches 53, Indels 3, Gaps 1;

```

```

Oy      223 COTGACCTCAGTGGATGATGTGAATTGTTCAATTCATGCAGGCCATCGTGAG 282
Db      207 CCTCAGCTCCTCATGTACTGACTGGAAFTCTTGCGCTTCTTCAACCCTGGGCATCATGCTGAG 148
Oy      283 CACTGTGAATCCAGAAGACGGGAACAACCTCCAAATGACCCCCTACCAACCAGTACATG--T 339
Db      147 CTACATCGCTCCAAAGAAAGCTGGAGACTCGAAGACCCATTCAAACGCTCATCATGAGTC 88
Oy      340 AGAGACTCGGCAAGAAAAGTCAAGAGCCAATC 373
Db      87 CGATGCTCGCAAGAGAAGAACAGGCTCATGTC 54

RESULT 9
US-09-864-761-17593/C
Sequence 17593, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharron G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
PRIOR FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
SEQ ID NO 17593
LENGTH: 390
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000168.1
OTHER INFORMATION: EXPRESSED IN HELLA, SIGNAL = 2.6
```

```
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 3.9
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.8
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.8
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 2.8
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.8
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 2.3
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3.2
; OTHER INFORMATION: EST_HUMAN HIT: AAT70188.1, EVALUE 6.30e-02
; OTHER INFORMATION: SWISSPROT HIT: P15382, EVALUE 4.00e-61
; OTHER INFORMATION: NT HIT: g11526222, EVALUE 0.00e+00
US-09-864-761-17593
```

```
Query Match 7.3%; Score 53.2; DB 10; Length 390;
Best Local Similarity 63.6%; Pred. No. 4.2e-06;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;
```

```
OY 223 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 282
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 209 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 150
OY 283 CACTGTGAATTCAGAGACGGGAACTCCATCATGACCCCTTACACCACTGATTTG--T 339
    | ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 149 CTACATCCGCTCCAGAACCTGGAGACCTCGAACGACCATTCACGCTCATCGAGTC 90
OY 340 AGAGACTGGCAGAAATACAGAGCCCAATC 373
    || ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 89 CGATGCTGTGCAAGAGAACGACGCTATGTC 56
```

```
RESULT 10
US-10-138-316-3
; Sequence 3, Application US/10138316
; Publication No. US20030054380A1
; GENERAL INFORMATION:
; APPLICANT: Keating, Mark T.
; APPLICANT: Sanguinetti, Michael C.
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MIK1 WHICH
; TITLE OF INVENTION: CAUSE ARRHYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING
; FILE REFERENCE: 2323-162
; CURRENT APPLICATION NUMBER: US/10/138,316
; PRIOR FILING DATE: 2002-05-06
; PRIOR APPLICATION NUMBER: 09/444,295
; PRIOR FILING DATE: 1999-11-22
; PRIOR APPLICATION NUMBER: 09/135,020
; PRIOR FILING DATE: 1998-08-17
; PRIOR APPLICATION NUMBER: 08/921,068
; PRIOR FILING DATE: 1997-08-29
; PRIOR APPLICATION NUMBER: 08/739,383
; PRIOR FILING DATE: 1996-10-29
; PRIOR APPLICATION NUMBER: 60/019,014
; PRIOR FILING DATE: 1995-12-22
; PRIOR APPLICATION NUMBER: 60/094,477
; PRIOR FILING DATE: 1998-07-29
; NUMBER OF SEQ ID NOS: 114
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 3
; LENGTH: 1703
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (193)..(579)
; FEATURE:
; NAME/KEY: misc
; LOCATION: (1)..(1703)
; OTHER INFORMATION: n may be any nucleotide
US-10-138-316-3
```

```
Query Match 7.3%; Score 53.2; DB 9; Length 1703;
```

```
Best Local Similarity 63.6%; Pred. No. 1e-05;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;
```

```
OY 223 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 282
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 324 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 383
OY 283 CACTGTGAATTCAGAGACGGGAACTCCATCATGACCCCTTACACCACTGATTTG--T 339
    | ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 384 CTACATCCGCTCCAGAACCTGGAGACCTCGAACGACCATTCACGCTCATCGAGTC 443
OY 340 AGAGACTGGCAGAAATACAGAGCCCAATC 373
    || ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 444 CGATGCTGTGCAAGAGAACGACGCTATGTC 477
```

```
RESULT 11
US-10-227-195A-1/c
; Sequence 1, Application US/10227195A
; Publication No. US20030077633A1
; GENERAL INFORMATION:
; APPLICANT: Cox, David
; APPLICANT: Arnold, Deana
; TITLE OF INVENTION: Haplotype structure of chromosome 21
; FILE REFERENCE: 103001
; CURRENT APPLICATION NUMBER: US/10/227,195A
; PRIOR FILING DATE: 2002-11-18
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 113604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7175, 7204, 36973, 66372, 76921, 81512, 88727
; OTHER INFORMATION: n = G or C
US-10-227-195A-1
```

```
Query Match 7.3%; Score 53.2; DB 9; Length 113604;
Best Local Similarity 63.6%; Pred. No. 0.00014;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;
```

```
OY 223 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 282
    ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 96499 CCTGTACCTCATGTGATGATTTGAATGTTCTTTTCATCATCGTGGCCATCGTGTGAG 96440
OY 283 CACTGTGAATTCAGAGACGGGAACTCCATCATGACCCCTTACACCACTGATTTG--T 339
    | ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 96439 CTACATCCGCTCCAGAACCTGGAGACCTCGAACGACCATTCACGCTCATCGAGTC 96380
OY 340 AGAGACTGGCAGAAATACAGAGCCCAATC 373
    || ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
DB 96379 CGATGCTGTGCAAGAGAACGACGCTATGTC 96346
```

```
RESULT 12
US-10-227-195A-2/c
; Sequence 2, Application US/10227195A
; Publication No. US20030077633A1
; GENERAL INFORMATION:
; APPLICANT: Cox, David
; APPLICANT: Arnold, Deana
; TITLE OF INVENTION: Haplotype structure of chromosome 21
; FILE REFERENCE: 103001
; CURRENT APPLICATION NUMBER: US/10/227,195A
; PRIOR FILING DATE: 2002-11-18
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 113604
; TYPE: DNA
; ORGANISM: Human
```



Query Match	6.3%	Score 46:	DB 10:	length 381:
Best Local Similarity	64.2%	Pred. No.	0.00069:	
Matches	86:	Conservative	0:	Mismatches 45: Indels 3: Gaps 1:

RESULT 15  
US-09-853-386-111  
; Sequence 111, Application US/09853386

```

: APPLICANT: Murphy, Evelyn
: APPLICANT: Bresnahan, Barry
: APPLICANT: Conneely, Orla
: APPLICANT: Fitzgerald, Oliver
: TITLE OF INVENTION: Therapeutic Approaches to Diseases by Suppression of the NURRR
: TITLE OF INVENTION: Subfamily of Nuclear Transcription Factors
: FILE REFERENCE: P01972051
: CURRENT APPLICATION NUMBER: US/09/853,386
: CURRENT FILING DATE: 2001-05-11
: PRIOR APPLICATION NUMBER: US 60/203645
: PRIOR FILING DATE: 2000-05-12
: NUMBER OF SEQ. ID NOS: 153
: SOFTWARE: PatentIn version 3.1
: SEQ. ID NO. 111
: LENGTH: 1146
: TYPE: DNA
: ORGANISM: HUMAN
: US-09-853-386-111

```

Query Match	4.9%	Score 36;	DB 10;	Length 1146;
Best Local Similarity	52.7%	Pred. No. 1.7;		
Matches 78;	Conservative 0;	Mismatches 70;	Indels 0;	Gaps 0;

188 GCCAAGTTGATGCTGAGACTTCTACTATGTCATCCTGTACCTCATGGTGATGATTGGA 247

Accession	Sequence	Position
Dd	GGCAAAAGCCCTGGGGGTGTACACGACTACATCTACACGAGGCCCCCATGATCTGTGTCCTG	738
Oy	ATGTTCTCTTTCATCAATCGTGGCCATCTGTGTGACACTGTGAATCCAAAGACGGAA	307
Dd	CTGATCAATTTTCATCTCTTCCTTTTAAACATCTGTCGACATCTCAATGACCAAGCTCCGGGCA	798
Oy	CACATCAATGACCCCTACCAACCAAGTACA	335
Dd	TTCACCAAGCTGTGAGACCATTCAGTACA	826

Search completed: May 21, 2003, 22:42:03  
Job time : 217.392 secs



GenCore version 5.1.4\_p5\_4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 20:14:00 ; Search time 42.5504 Seconds  
(without alignments)  
575.799 Million cell updates/sec

Title: US-09-550-163-1

Sequence: 1 caaatccagaagaatccgt.....atgaataataagccaattt 732

Scoring table:

IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 segs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued Patents, NA:  
1: /cgn2\_6/ptodata/1/lna/5A.COMB.seq:\*  
2: /cgn2\_6/ptodata/1/lna/5B.COMB.seq:\*  
3: /cgn2\_6/ptodata/1/lna/6A.COMB.seq:\*  
4: /cgn2\_6/ptodata/1/lna/6B.COMB.seq:\*  
5: /cgn2\_6/ptodata/1/lna/PCUTS.COMB.seq:\*  
6: /cgn2\_6/ptodata/1/lna/Backfiles1.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	53.2	7.3	398	1	US-08-118-101A-5
2	53.2	7.3	436	4	US-09-679-185-1
3	53.2	7.3	1703	3	US-09-135-021-77
4	53.2	7.3	1703	4	US-09-135-020-3
5	53.2	7.3	1703	4	US-09-135-010A-3
6	53.2	7.3	1703	4	US-09-444-871-3
7	53.2	7.3	1703	4	US-09-597-735-3
8	53.2	7.3	1703	4	US-09-444-295-3
9	53.2	7.3	1703	4	US-09-597-733-3
10	51.6	7.0	436	4	US-09-679-185-3
11	36.6	5.0	2652	1	US-08-318-831-1
12	36.2	4.9	7218	1	US-08-232-463-14
13	36	4.9	1380	1	US-08-110-286A-1
14	36	4.9	1495	4	US-08-482-746-1
15	36	4.9	1582	4	US-08-482-746-14
16	33.6	4.6	645	3	US-09-069-896-2
17	33.6	4.6	645	4	US-09-471-468-2
18	33	4.5	606	4	US-09-328-111-133
19	33	4.5	2912	4	US-09-307-143-3
20	32.4	4.4	837	4	US-08-998-416-303
21	32.4	4.4	4659	4	US-09-221-017B-823
22	32.2	4.4	43676	3	US-09-356-952-12
23	32	4.4	246240	2	US-08-724-394A-20
24	32	4.4	246240	2	US-08-724-394A-21
25	32	4.4	246240	2	US-08-724-394A-22
26	31.4	4.3	566	4	US-09-221-017B-919
27	30.6	4.2	744	3	US-08-969-644-17

28	30.6	4.2	744	3	US-08-444-189-17	Sequence 17, Appl
29	30.6	4.2	744	4	US-08-468-544-17	Sequence 17, Appl
30	30.6	4.2	2929	4	US-09-705-299-10	Sequence 10, Appl
31	30.6	4.2	6822	4	US-09-426-998-3	Sequence 3, Appl
32	30.6	4.2	7502	3	US-08-969-644-6	Sequence 6, Appl
33	30.6	4.2	7502	3	US-08-444-189-6	Sequence 6, Appl
34	30.6	4.2	7502	4	US-08-468-544-6	Sequence 6, Appl
35	30.6	4.2	7741	4	US-09-426-998-4	Sequence 4, Appl
36	30.6	4.2	98844	4	US-09-791-211-10	Sequence 10, Appl
37	30.4	4.2	28473	4	US-08-961-527-83	Sequence 83, Appl
38	30.2	4.1	28001	4	US-09-819-993-3	Sequence 3, Appl
39	30.2	4.1	33000	4	US-09-215-694-18	Sequence 18, Appl
40	30	4.1	1001	4	US-09-641-638-198	Sequence 198, App
41	29.8	4.1	2266	2	US-08-724-394A-18	Sequence 18, Appl
42	29.8	4.1	72604	4	US-09-268-992-7	Sequence 7, Appl
43	29.8	4.1	72604	4	US-09-657-474-7	Sequence 7, Appl
44	29.6	4.0	462	3	US-08-863-813A-33	Sequence 33, Appl
45	29.6	4.0	3273	6	5516630-1	Patent No. 5516630

# ALIGNMENTS

RESULT 1  
US-08-118-101A-5

Sequence 5, Application US/08118101A  
Patent No. 5620892

GENERAL INFORMATION:

APPLICANT: Kutz, Stephen E.  
APPLICANT: Knickerbocker, Aron M.

TITLE OF INVENTION: A STRAIN OF SACCHAROMYCES CEREVISIAE  
TITLE OF INVENTION: EXPRESSING THE GENE ENCODING POTASSIUM TRANSPORTER MINK

NUMBER OF SEQUENCES: 16  
CORRESPONDENCE ADDRESSES:

ADDRESSEE: Burton Rodney  
STREET: P O. Box 4000

CITY: Princeton  
STATE: New Jersey

COUNTRY: U.S.A.  
ZIP: 08543-4000

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/118,101A  
FILING DATE:

CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:

NAME: Gaul, Timothy J.  
REGISTRATION NUMBER: 33,111

REFERENCE/DOCKET NUMBER: DC27  
TELECOMMUNICATION INFORMATION:

TELEPHONE: (609) 252-5901  
TELEFAX: (609) 252-4526

INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:

LENGTH: 398 base pairs  
TYPE: nucleic acid

STRANDEDNESS: single  
TOPOLOGY: linear

MOLECULE TYPE: cDNA  
FEATURE:

NAME/KEY: CDS  
LOCATION: 1..398

US-08-118-101A-5

Query Match 7.3%; Score 53.2; DB 1; Length 398;  
Best Local Similarity 63.6%; Pred. No. 3.4e-07;  
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;











Query Match 4.9%; Score 36.2; DB 1; Length 7218;  
 Best Local Similarity 2.3%; Pred. No. 0.4;  
 Matches 8; Conservative 191; Mismatches 144; Indels 0; Gaps 0;

```

QY 277 GGTAGACACTGTGAATCCAGAGACGGGAACTCCATGACCCCTACCAACAGATACAT 336
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1398 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1339
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 337 TGTAGAGACTGCGAGAAAGTACAGACCAATCTGATCTAGAGATGGAAGGC 396
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1338 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1279
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 397 CACCATCATGAGAACTGTGGGCTGGTTCAAAATGTCCTCCCTGATAGGAGAAA 456
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1278 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1219
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 457 GGCACCAAGCTAACATGTGACGTCAGACATGAAAGAGATGCCAGGAGCAATC 516
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1218 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1159
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 517 CAATATGCTTTGCTAGAGAAAGTGAATGCTCTTGTGTGATTTTCATGGA 576
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1158 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1099
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 577 GATTATGCTGTGGCAATAAGATAGATGACATTTCAATCTC 619
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
DB 1098 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1056
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
  
```

## RESULT 13

```

US-08-110-2864-1
; Sequence 1, Application US/08110286A
; Patent No. 5728545
; GENERAL INFORMATION:
; APPLICANT: Perrin, Marilyn H.
; APPLICANT: Chen, Ruoping
; APPLICANT: Lewis, Kathy A.
; APPLICANT: Vale Jr., Wylie W.
; APPLICANT: Donaldson, Cynthia J.
; TITLE OF INVENTION: CLONING AND RECOMBINANT PRODUCTION OF
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pretty, Schroeder, Brueggemann & Clark
; STREET: 444 South Flower Street, Suite 2000
; CITY: Los Angeles
; STATE: CA
; COUNTRY: USA
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/110,286A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/079,320
; FILING DATE: 18-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Reiter, Stephen E.
; REGISTRATION NUMBER: 31,192
; REFERENCE/DOCKET NUMBER: P41 9439
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-546-4737
; TELEFAX: 619-546-9392
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1380 base pairs
; TYPE: nucleic acid
  
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STRANDEDNESS: both  
 TOPOLOGY: both  
 MOLECULE TYPE: cDNA  
 FEATURE:

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; NAME/KEY: CDS
; LOCATION: 82..1329
; OTHER INFORMATION: /product="HUMAN PITUITARY
; OTHER INFORMATION: CRF-RECEPTOR
; OTHER INFORMATION: /note="This sequence is encoded by clone
; OTHER INFORMATION: "CRF-R1".
US-08-110-2864-1
  
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## RESULT 14

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US-08-482-746-1
; Sequence 1, Application US/08482746B
; Patent No. 6399315
; GENERAL INFORMATION:
; APPLICANT: Perrin, Marilyn H.
; APPLICANT: Chen, Ruoping
; APPLICANT: Lewis, Kathy A.
; APPLICANT: Vale Jr., Wylie W.
; APPLICANT: Donaldson, Cynthia J.
; APPLICANT: Savchenko, Paul
; TITLE OF INVENTION: Cloning and Recombinant Production of
; NUMBER OF SEQUENCES: 6
; FILE REFERENCE: P41-90002
; CURRENT APPLICATION NUMBER: US/08/482,746B
; EARLIER FILING DATE: 1995-06-07
; EARLIER APPLICATION NUMBER: US 08/374,009
; EARLIER FILING DATE: 1995-01-17
; EARLIER APPLICATION NUMBER: US 08/353,537
; EARLIER FILING DATE: 1994-12-09
; EARLIER APPLICATION NUMBER: PCT/US94/05908
; EARLIER FILING DATE: 1994-05-25
; EARLIER APPLICATION NUMBER: US 08/110,286
; EARLIER FILING DATE: 1993-08-23
; EARLIER APPLICATION NUMBER: US 08/079,320
; EARLIER FILING DATE: 1993-06-18
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 1495
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (82)...(1326)
; OTHER INFORMATION: /product="Human pituitary CRF-receptor"
; OTHER INFORMATION: /note="this sequence is encoded by clone
; OTHER INFORMATION: "CRF-R1".
US-08-482-746-1
  
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 Best Local Similarity 52.7%; Pred. No. 0.19;  
 Matches 78; Conservative 0; Mismatches 70; Indels 0; Gaps 0;





GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:16:44 ; Search time 178.18 Seconds

(without alignments)  
9251.680 Million cell updates/sec

Title: US-09-550-163-1

Perfect score: 732  
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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	732	100.0	732	21 AAC64071	Human potassium ch
2	732	100.0	732	24 AAD35170	Human KCNE2 wild t
3	730.4	99.8	732	21 AAC64083	Human potassium ch
4	730.4	99.8	732	21 AAC64084	Human potassium ch
5	730.4	99.8	732	21 AAC64085	Human potassium ch
6	730.4	99.8	732	21 AAC64086	Human potassium ch
7	730.4	99.8	732	24 AAK86573	cDNA encoding huma
8	730.4	99.8	732	24 AAD35169	Human KCNE2 mutant
9	730.4	99.8	732	24 AAD35171	Human KCNE2 mutant

10	730.4	99.8	732	24 AAD35172	Human KCNE2 mutant
11	730.4	99.8	732	24 AAD35173	Human KCNE2 mutant
12	553.8	75.7	600	22 ABA09192	Human MIRP1 homolo
13	553.8	75.7	600	22 AAK52645	Human polynucleotl
14	553.8	75.7	655	22 AAK51661	Human polynucleotl
15	466	63.7	471	22 AAF80269	Nucleotide sequenc
16	372	50.8	372	22 AA124432	Probe #14365 for g
17	372	50.8	372	22 AA109965	Probe #9956 used t
18	372	50.8	372	22 AAS00245	Human potassium ch
19	368	50.3	450	22 ABA44797	Human breast cell
20	368	50.3	450	22 ABA45252	Human foetal liver
21	368	50.3	450	22 ABA24997	Probe #3463 for ge
22	368	50.3	450	22 AAK03508	Human brain expres
23	368	50.3	450	22 AAK28962	Human bone marrow
24	368	50.3	450	22 AAT13549	Probe #3482 for ge
25	368	50.3	450	22 AAT14911	Probe #3597 used t
26	368	50.3	450	22 AAI03438	Probe #3429 used t
27	368	50.3	450	22 ABS03496	Human genome-deriv
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29	312	42.6	312	22 ABA67856	Human foetal liver
30	312	42.6	312	22 ABA34913	Probe #13379 for g
31	312	42.6	312	22 AAK16264	Human brain expres
32	312	42.6	312	22 AAK42008	Human bone marrow
33	312	42.6	312	22 AAI22773	Probe #12706 for g
34	312	42.6	312	22 AAI48075	Probe #16761 used
35	312	42.6	312	22 AAI08446	Probe #8437 used t
36	312	42.6	312	24 ABS16039	Human genome-deriv
37	306	41.8	471	22 AAI15256	Probe #5189 for ge
38	306	41.8	471	22 AAI04990	Probe #4981 used t
39	277.4	37.9	468	21 AAC64072	Rat potassium chan
40	266.4	36.4	372	22 AAS00246	Rat potassium chan
41	60	8.2	60	24 ABA37461	Human spliced tran
42	53.8	7.3	65	24 ABA29241	Rat spliced transc
43	53.2	7.3	231	22 ABA68478	Human foetal liver
44	53.2	7.3	231	22 ABA35463	Human KCNE2 for g
45	53.2	7.3	231	22 AAK16847	Human brain expres

#### ALIGNMENTS

RESULT 1	
AAC64071	
ID AAC64071 standard; cDNA: 732 BP.	
XX AAC64071;	
DT 19-FEB-2001 (first entry)	
XX	
AC	Human potassium channel protein KCNE2 (MIRP1) cDNA, SEQ ID NO:1.
XX	
DE	Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;
XX	MIRK-related; long QT syndrome; cardiac arrhythmia;
KW	drug screening; knock out mouse; transgenic animal; ion channel disorder;
KW	fast delayed rectifier potassium channel; anti-KCNE2 antibody;
KW	HENG; ss.
XX	
OS	Homo sapiens.
XX	
PN	WO200063434-A1.
XX	
PD	26-OCT-2000.
XX	
PF	14-APR-2000; 2000WO-US10004.
XX	
PR	15-APR-1999; 99US-0129404.
XX	
PA	(UTAH ) UNIV UTAH RES FOUND.
PA	(UTAH ) UNIV YALE.
XX	
PI	Abbolt GW, Sesti F, Splawski I, Keating MT, Goldstein SAN;
XX	WPI, 2000-672747/65.

DR P-PSDB; AAB29585.  
 XX Novel nucleic acids encoding M1RP1, M1RP2 and M1RP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 XX  
 PS Claim 1; Page 118-119; 132pp; English.  
 XX  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (M1RP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (M1RP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (M1RP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-Kr), mutations in which are associated with long  
 CC QT syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents cDNA encoding human KCNE2 (M1RP1).  
 XX  
 SO Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other;  
 Query Match 100.0%; Score 732; DB 21; Length 732;  
 Best Local Similarity 100.0%; Pred. No. 3.2e-203;  
 Matches 732; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 .  
 QY 1 CAATTCAGAAAGATGCGTTTCTTACCTTGTGCGCTATTTATTTAAATGCA 60  
 DB 1 CAATTCAGAAAGATGCGTTTCTTACCTTGTGCGCTATTTATTTAAATGCA 60  
 QY 61 GCAGAGGAGGAGCATGCTCTATTATTCAAATTTACACAGACGCTGAGACGCTCCG 120  
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 DB 301 ACGGGAACATCCCATATGACCCCTACACAGACATGATGAGAGACTGGAGAAAAGTA 360  
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 DB 361 CAAGAGCAATCTTGAATGATAGAGAAATGGAAGGCCACATCCATAGAAACATTTGGTGC 420  
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 DB 661 AGACCTCTTTTACTTTCCGGGCAAGTGAATGTCATTTTATCAATATCATATGAAAT 720  
 QY 721 AAGCCAAATTT 732  
 DB 721 AAGCCAAATTT 732

## RESULT 2

ID AAD35170 standard; DNA; 732 BP.

XX AAD35170;

XX 25-JUL-2002 (first entry)

XX Human KCNE2 wild type DNA.

XX Human; Min-K related ion channel protein; M1RP1; ion channel disorder;

KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; gene; ds.

XX Homo sapiens.

XX Key Location/Qualifiers

FT CDS 74..445 /tag= a /product= "Human M1RP1 protein"

PN W0200222875-A2.

XX 21-MAR-2002.

XX 11-SEP-2001; 2001WO-US28332.

XX 11-SEP-2000; 2000US-231571P.

XX (UYTA ) UNIV YALE.

XX Goldstein SAN;

DR M1P1: 2002-362360/39.

DR P-PSDB; AAE22095.

XX Novel gene encoding Min-K related ion channel protein subunit and

PT polymorphisms in this gene associated with antibiotic-induced long QT

PT syndrome -

PS Claim 9; Page 43; 49pp; English.

XX The present invention relates to novel KCNE2 genes encoding Min-K related

CC (M1RP) 1 ion channel proteins and polymorphisms in these genes that are

CC associated with ion channel disorders including antibiotic-induced long

CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,

CC 57 or 116 of M1RP1 polypeptide, or a mutation at a nucleotide position

CC encoding the amino acid positions is useful for diagnosing the presence

CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods

CC are useful in the development of new drug therapies which selectively

CC target one or more KCNE2 polymorphisms that are associated with cardiac

CC arrhythmias. The present sequence is human KCNE2 wild type DNA.

SO Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other;

Query Match 100.0%; Score 732; DB 24; Length 732;

Best Local Similarity 100.0%; Pred. No. 3.2e-203;

Matches 732; Conservative 0; Mismatches 0; Indels 0; Gaps 0;



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 DB 421 GGGTGGGTTCAAAATGTCCCTGATATAGGAGAAAGGACCAAGCTAACATCTGACGTC 480  
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 DB 481 CAGACATGAAAGATGCCAGTGGCCAGAGCAATCCAAATTTGCTTCTTGAAGAAA 540  
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 DB 661 AGACCTTTTACTTCCGCGCAAGTGAATTCATTTTAAATCAATGATGAAT 720  
 QY 721 AAAGCCAAATTT 732  
 DB 721 AAAGCCAAATTT 732  
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 AAC64083  
 ID AAC64083 standard; DNA; 732 BP.  
 XX AAC64083;  
 XX 19-FEB-2001 (first entry)  
 DE Human potassium channel protein KCNE2 (MIRP1) Q9E mutant DNA.  
 XX  
 XX Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
 KM Mink-related; long QT syndrome; cardiac arrhythmia;  
 KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KM HERG; mutant; ds.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 PN WO200063434-A1.

XX 26-OCT-2000.  
 PD 14-APR-2000; 2000WO-US10004.  
 XX 15-APR-1999; 99US-0129404.  
 PF (UTAH ) UNIV UTAH RES FOUND.  
 XX (UTAH ) UNIV YALE.  
 PA Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAN;  
 PI WPI: 2000-672747/65.  
 DR P-FSDB; AAB29593.  
 XX Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 XX  
 PS Claim 56; Page -; 132pp; English.  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-KR), mutations in which are associated with long QT  
 CC syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents DNA encoding a mutant human KCNE2  
 CC (MIRP1) specifically claimed for use in diagnostic and drug screening  
 CC methods of the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
 CC 118-119.  
 SQ Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other;  
 Query Match 99.8%; Score 730.4; DB 21; Length 732;  
 Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
 Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
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 DB 1 CAAATCCAGAAAAAGATCGGTTTCTACCTGTTGCGCCTATTTTATTTAAATTGCA 60  
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 DB 61 GCAGAGGAGAGATGCTACTTATTCATATTCACACAGAGCGTGAAGAGCTCTCCG 120  
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 DB 121 AAGGATTTTATTTACTTATATGACAAATTTGGCGCCAGAAACACACAGCTGACCAAGAGC 180  
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 DB 181 CCTCCAGCCAAAGTTGATGATGAGAACTTCTACTATGTCATCCCTGCTACTCATGCTGAT 240  
 QY 241 GATTGGAATGTTCTCTTTTCATCATCGTGGCCATCCCTGGTGGACACTGTGAATCCCAAGAG 300  
 DB 241 GATTGGAATGTTCTCTTTTCATCATCGTGGCCATCCCTGGTGGACACTGTGAATCCCAAGAG 300

QY	301	ACGGGAACACCCCAATGACCCCTCCACCCACGTAATGTAGAGACGTGGCAAGAAAGTA	360
Db	301	ACGGGAACACCCCAATGACCCCTCCACCCACGTAATGTAGAGACGTGGCAAGAAAGTA	360
QY	361	CAAGAGCCCAATCTTGAATCTAGAAGAAATCGAAGGCCACCATCCATGAGAAATTTGGTGC	420
Db	361	CAAGAGCCCAATCTTGAATCTAGAAGAAATCGAAGGCCACCATCCATGAGAAATTTGGTGC	420
QY	421	GGCTGGGTTCAAAATGTCCCTCTATTAAGGAGAAAGGACCAAGCTAACATCTGACGTC	480
Db	421	GGCTGGGTTCAAAATGTCCCTCTATTAAGGAGAAAGGACCAAGCTAACATCTGACGTC	480
QY	481	CAGACATGAGAGATGTCAGTGCACGACGAGCAATCCAAATTCCTTGGCTTAGAAGAA	540
Db	481	CAGACATGAGAGATGTCAGTGCACGACGAGCAATCCAAATTCCTTGGCTTAGAAGAA	540
QY	541	GTCAGTTCCTTCCTTGGTGTGAGAAATTTTCATGAGAGATTAATGTGGTGGCCAAATAAGA	600
Db	541	GTCAGTTCCTTCCTTGGTGTGAGAAATTTTCATGAGAGATTAATGTGGTGGCCAAATAAGA	600
QY	601	TAGATGACATTTCAATTCAGATGATTAATGCTTGTGAGACATATTTTGTGCTGA	660
Db	601	TAGATGACATTTCAATTCAGATGATTAATGCTTGTGAGACATATTTTGTGCTGA	660
QY	661	AGACCTCTTTTACTTTCGCGGCAAGTGAATGTCATTTTAAATCAATCAATGATGAAT	720
Db	661	AGACCTCTTTTACTTTCGCGGCAAGTGAATGTCATTTTAAATCAATCAATGATGAAT	720
QY	721	AAAGCCAAATTT 732	
Db	721	AAAGCCAAATTT 732	
RESULT 4			
AAC64084			
ID	AAC64084 standard; DNA: 732 BP.		
AC	AAC64084:		
AT	19-FEB-2001 (first entry)		
DE	Human potassium channel protein KCNE2 (MiRP1) M54T mutant DNA.		
KW	Human: KCNE2: MiRP1: potassium channel protein; KCNE1-related;		
KM	Mink-related; long QT syndrome; cardiac arrhythmia;		
KW	drug screening; knockout mouse; transgenic animal; ion channel disorder;		
KM	fast delayed rectifier potassium channel; anti-KCNE2 antibody;		
KW	HERG; mutant; ds.		
OS	Homo sapiens.		
OS	Synthetic.		
PN	WO200063434-A1.		
XX	26-OCT-2000.		
XX	14-APR-2000; 2000MO-US10004.		
XX	15-APR-1999; 99US-0129404.		
XX	(UTAH ) UNIV UTAH RES FOUND.		
XX	(UYVA ) UNIV YALE.		
PI	Abbott GW, Seethi F, Splawski I, Keating MT, Goldstein SAN;		
DR	WPI; 2000-672747/65.		
XX	P-PSDB; AAB29594.		
PT	Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for		
PT	diagnosing and treating ion channel disorders, especially long QT		
XX	syndrome -		
XX	Claim 56; Page -; 132pp; English.		

The invention relates to novel ion channel proteins related to KCNE1 (Mink) and to nucleic acids encoding them. The proteins of the invention are human and rat KCNE2 (MRP2; AAB29585 and AAB29586, respectively); human and mouse KCNE3 (MRP3; AAB29587 and AAB29588, respectively); and human and mouse KCNE4 (MRP3; AAB29589 and AAB29590, respectively). The cDNAs encoding these proteins are given in AAC64071-AAC64076. KCNE2, along with HENK, forms cardiac fast delayed rectifier potassium channels (I-KR), mutations in which are associated with long QT syndrome. The invention also relates to methods of diagnosing long QT syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic nonhuman animals comprising a heterologous ion channel protein gene of the invention, a transgenic animal comprising human KCNE2 and HENK DNA, and methods of and screening drugs for treating long QT syndrome using KCNE2 proteins (including mutants), nucleic acids encoding them and antibodies against KCNE2 proteins. The methods, antibodies, nucleic acids, and proteins may be used for diagnosing or treating ion channel disorders, especially long QT syndrome. Transgenic animals comprising KCNE2 and HENK are useful for testing anti-long QT syndrome drugs.

The present sequence represents DNA encoding a mutant human KCNE2 (MRP1) specifically claimed for use in diagnostic and drug screening methods of the invention.

Note: The present sequence is not shown in the specification, but is derived from the wild-type human KCNE2 cDNA sequence shown on page 118-119.

Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match	99.8%	Score 730.4	DB 21	Length 732
Best Local Similarity	99.9%	Pred. No. 9.2e-203		
Matches 731	Conservative	0	Mismatches 1	Indels 0
				Gaps 0
QY 1	CAATCCAGAAAAGATCCGTTTCTTAACCTGTTCGGCTATTATATATTAATTAATGA	60		
Db 1	CAATCCAGAAAAGATCCGTTTCTTAACCTGTTCGGCTATTATATATTAATTAATGA	60		
QY 61	GCAGGAGGGAAGCATGTCTAATTATCCATTTCACACAGACGCTGGACAGCGTTCCG	120		
Db 61	GCAGGAGGGAAGCATGTCTAATTATCCAAATTTCAACAGACGCTGGACAGCGTTCCG	120		
QY 121	AAGGATTTTATTAATCTTAATGATCAATTTGGGGCCAGAACCAACAGCTGGACAAAGGC	180		
Db 121	AAGGATTTTATTAATCTTAATGATCAATTTGGGGCCAGAACCAACAGCTGGACAAAGGC	180		
QY 181	CCTCCAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGTAT	240		
Db 181	CCTCCAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGTAT	240		
QY 241	GATTGGAAATTTCTTTTATCATCTGTGGCCATCCTGGTGAGCACTGTGAAATCCAGAG	300		
Db 241	GATTGGAAATTTCTTTTATCATCTGTGGCCATCCTGGTGAGCACTGTGAAATCCAGAG	300		
QY 301	ACGGGAACATCTCAATAGACCCCTACACACAGTATTTAAGAGACTGGCAAAAAGTA	360		
Db 301	ACGGGAACATCTCAATAGACCCCTACACACAGTATTTAAGAGACTGGCAAAAAGTA	360		
QY 361	CAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACCATCATGAGAACTTGGTGC	420		
Db 361	CAGAGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACCATCATGAGAACTTGGTGC	420		
QY 421	GGCTGGGTTCAAAATGTCCCTCGTAATAGGGGAAGGACACCAAGCTPAACATCTGCAGCTC	480		
Db 421	GGCTGGGTTCAAAATGTCCCTCGTAATAGGGGAAGGACACCAAGCTPAACATCTGCAGCTC	480		
QY 481	CACAGCATGAAGATGCCAGTGCACACGAGGCAAAATCTGCTTTGTGTAGAAAGAA	540		
Db 481	CACAGCATGAAGATGCCAGTGCACACGAGGCAAAATCTGCTTTGTGTAGAAAGAA	540		
QY 541	GTGAGTTCTTGTCTTTGTAGAAATTTCTAGAGATATGTGTGTGGCCAAATAAGA	600		
Db 541	GTGAGTTCTTGTCTTTGTAGAAATTTCTAGAGATATGTGTGTGGCCAAATAAGA	600		

QY 601 TAGATGACATTTCAATCTCAGATTTATGCTTGTGTCGAGCAAAATATTTTGTGCTGA 660  
|||||  
DB 601 TAGATGACATTTCAATCTCAGATTTATGCTTGTGTCGAGCAAAATATTTTGTGCTGA 660  
QY 661 AGACCTCTTTACTTTCGCGGCAAGTGAATGCTATTTAACTAATATCAATGATGATAAT 720  
|||||  
DB 661 AGACCTCTTTACTTTCGCGGCAAGTGAATGCTATTTAACTAATATCAATGATGATAAT 720  
QY 721 AAAGCCAAATTT 732  
|||||  
DB 721 AAAGCCAAATTT 732  
RESULT 5  
AAC64085  
ID AAC64085 standard; DNA; 732 BP.  
XX  
AC AAC64085;  
XX  
DT 19-FEB-2001 (first entry)  
XX  
DE Human potassium channel protein KCNE2 (MIRP1) I57T mutant DNA.  
XX  
KM Human: KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KM Mink-related; long QT syndrome; cardiac arrhythmia;  
KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KM HERG; mutant; ds.  
XX  
OS Homo sapiens.  
OS Synthetic.  
XX  
PN WO20063434-A1.  
XX  
PD 26-OCT-2000.  
XX  
PF 14-APR-2000; 2000WO-US10004.  
XX  
PR 15-APR-1999; 99US-0129404.  
XX  
PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYVA ) UNIV YALE.  
XX  
PI Abdoct GW, Sestl F, Splawski I, Keating WT, Goldstein SAN;  
DR WPI; 2000-672147/65.  
DR P-PSDB; AAB29595.  
XX  
PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
PT diagnosing and treating ion channel disorders, especially long QT  
PT syndrome -  
XX  
PS Claim 56; Page -; 132pp; English.  
XX  
CC The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-KR), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.

CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (MIRP1) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.  
CC Note: The present sequence is not shown in the specification, but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.  
XX  
SO Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;  
QY  
Query Match 99.8%; Score 730.4; DB 21; Length 732;  
Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCCGCTATTTATTTAATTTGCA 60  
DB 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCCGCTATTTATTTAATTTGCA 60  
QY 61 GCAGAGGGAAGCATGTCTACTTTATCCAAATTTACACACAGCGTGAAGACGTTCCG 120  
DB 61 GCAGAGGGAAGCATGTCTACTTTATCCAAATTTACACACAGCGTGAAGACGTTCCG 120  
QY 121 AAGGATTTTATTTACTTATATGACAAATGGCGCAGACACACAGCTGACGAAGGCG 180  
DB 121 AAGGATTTTATTTACTTATATGACAAATGGCGCAGACACACAGCTGACGAAGGCG 180  
QY 181 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCTATCCTGTACCTCATGTGAT 240  
DB 181 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCTATCCTGTACCTCATGTGAT 240  
QY 241 GATTGGAATGTTCTCTTTCATCATCGTGGCCATCCTGGTGAGCACTGTGAATCCAAAG 300  
DB 241 GACTGGAATGTTCTCTTTCATCATCGTGGCCATCCTGGTGAGCACTGTGAATCCAAAG 300  
QY 301 ACGGGAACATCTCAATGACCCCTACACACAGTACATGTAAGGACGCGAGAAAAGTA 360  
DB 301 ACGGGAACATCTCAATGACCCCTACACACAGTACATGTAAGGACGCGAGAAAAGTA 360  
QY 361 CAAGAGCCAAATCTGAAATCTAGAAGATCGAAGGCCACATCCATGAGAAATTTGGTGC 420  
DB 361 CAAGAGCCAAATCTGAAATCTAGAAGATCGAAGGCCACATCCATGAGAAATTTGGTGC 420  
QY 421 GCGTGGGTCAAAATGTCCTCCCTGATTAAGGAGAAAGCCAAAGCTTAACATCTGACGTC 480  
DB 421 GCGTGGGTCAAAATGTCCTCCCTGATTAAGGAGAAAGCCAAAGCTTAACATCTGACGTC 480  
QY 481 CAGACATGAAGAGATGCCAGTGCACGAGGCAAAATTCAAATTTGCTTGTAGAGAAA 540  
DB 481 CAGACATGAAGAGATGCCAGTGCACGAGGCAAAATTCAAATTTGCTTGTAGAGAAA 540  
QY 541 GTGAGTTCCTTGTCTTGTGAGAAATTTTCATGAGATTTGTTGGCCAAATAAGA 600  
DB 541 GTGAGTTCCTTGTCTTGTGAGAAATTTTCATGAGATTTGTTGGCCAAATAAGA 600  
QY 601 TAGATGACATTTCAATCTCAGATTTATGCTTGTGTCGAGCAAAATATTTTGTGCTGA 660  
DB 601 TAGATGACATTTCAATCTCAGATTTATGCTTGTGTCGAGCAAAATATTTTGTGCTGA 660  
QY 661 AGACCTCTTTACTTTCGCGGCAAGTGAATGCTATTTAACTAATATCAATGATGATAAT 720  
DB 661 AGACCTCTTTACTTTCGCGGCAAGTGAATGCTATTTAACTAATATCAATGATGATAAT 720  
QY 721 AAAGCCAAATTT 732  
DB 721 AAAGCCAAATTT 732  
RESULT 6  
AAC64086  
ID AAC64086 standard; DNA; 732 BP.  
XX  
AC AAC64086;  
XX  
DT 19-FEB-2001 (first entry)

XX Human potassium channel protein KCNE2 (MiRP1) T8A mutant DNA.  
 XX  
 XX Human; KCNE2; MiRP1; potassium channel protein; KCNE1-related;  
 KW Mink-related; long QT syndrome; cardiac arrhythmia;  
 KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KW HERG; mutant; ds.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 PN W0200063434-A1.  
 XX  
 PD 26-OCT-2000.  
 XX  
 PF 14-APR-2000; 2000MO-US10004.  
 XX  
 PR 15-APR-1999; 99US-0129404.  
 XX  
 PA (UTAH ) UNIV UTAH RES FOUND.  
 PA (UTIA ) UNIV IALE.  
 PI Abbott GW, Seatl F, Splawski I, Keating MT, Goldstein SAN;  
 PI WPI; 2000-672747/65.  
 DR P-PSDB; AAB29596.  
 XX  
 PT Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 PS  
 PS Claim 56; Page -; 132pp; English.  
 XX  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MiRP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (MiRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MiRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAB64071-  
 CC AAB64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-Kr), mutations in which are associated with long  
 CC QT syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents DNA encoding a mutant human KCNE2  
 CC (MiRP1) specifically claimed for use in diagnostic and drug screening  
 CC methods of the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
 CC 118-119.  
 XX  
 XX  
 SQ Sequence 732 BP; 220 A; 152 C; 158 G; 202 T; 0 other;

Query Match 99.8%; Score 730.4; DB 21; Length 732;  
 Best Local Similarity 99.8%; Pred. No. 9,2e-203;

Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAATCCAGAAAAGATCGTTTCTTACCTGTGCGCATTTTATTTAAATGCA 60  
 DB 1 CAATCCAGAAAAGATCGTTTCTTACCTGTGCGCATTTTATTTAAATGCA 60  
 QY 61 GCAGGAGGAGGAGCATCTTATTCATTTACACAGAGCGTGAAGACGCTCCG 120  
 DB 61 GCAGGAGGAGGAGCATCTTATTCATTTACATTTCCGACAGAGCGTGAAGACGCTCCG 120

QY 121 AAGGATTTTATTACTTATATGCAATTTGGCCGCAACACAGACGTGAGCAAGGC 180  
 DB 121 AAGGATTTTATTACTTATATGCAATTTGGCCGCAACACAGACGTGAGCAAGGC 180  
 QY 181 CTTCCAAAGCCAAAGTTGATGCTGAGAACTTCTATGTCATCTCTTACCTCATGTTGAT 240  
 DB 181 CTTCCAAAGCCAAAGTTGATGCTGAGAACTTCTATGTCATCTCTTACCTCATGTTGAT 240  
 QY 241 GATTGGAATGTTCTTTTCATTCATGTCGACCATCTGTTGAGCAGTGTAAATCCAAAG 300  
 DB 241 GATTGGAATGTTCTTTTCATTCATGTCGACCATCTGTTGAGCAGTGTAAATCCAAAG 300  
 QY 301 ACGGGAACCTCCCAATGACCCCTACGACCAAGTACATTTGAGAGACTGGCAGAAAAGTA 360  
 DB 301 ACGGGAACCTCCCAATGACCCCTACGACCAAGTACATTTGAGAGACTGGCAGAAAAGTA 360  
 QY 361 CAAGAGCCAAATCTTGAATCTAGAAATTCGAAAGCCACCATCTCATGAGAACATTTGTC 420  
 DB 361 CAAGAGCCAAATCTTGAATCTAGAAATTCGAAAGCCACCATCTCATGAGAACATTTGTC 420  
 QY 421 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGCACCAGCTACATCTGACGTC 480  
 DB 421 GGCTGGGTTCAAAATGTCCCCCTGATTAAGGAGAAAGCACCAGCTACATCTGACGTC 480  
 QY 481 CAGACATGAAGAGATGCCAGTGCACGAGCAAAATCCAAATGTCTTTGTTAGAGAA 540  
 DB 481 CAGACATGAAGAGATGCCAGTGCACGAGCAAAATCCAAATGTCTTTGTTAGAGAA 540  
 QY 541 GTGAGTCCCTGCTCTTGTGAGAAATTTCAATGAGATTAATGTTGGTGGCCAAATTAAGA 600  
 DB 541 GTGAGTCCCTGCTCTTGTGAGAAATTTCAATGAGATTAATGTTGGTGGCCAAATTAAGA 600  
 QY 601 TAGATGACATTTCAATCTCAGTATTTATCTGCTTGTGAGCAATTTTGTGCTGA 660  
 DB 601 TAGATGACATTTCAATCTCAGTATTTATGCTTGTGAGCAATTTTGTGCTGA 660  
 QY 661 AGACCTCTTTTACTTCCGGGCAAGTGAATGCAATTTTAAATATCAATGATGAAT 720  
 DB 661 AGACCTCTTTTACTTCCGGGCAAGTGAATGCAATTTTAAATATCAATGATGAAT 720  
 QY 721 AAAGCCAAATTT 732  
 DB 721 AAAGCCAAATTT 732

RESULT 7  
 ABR6573  
 ID ABR6573 strand: DNA; 732 BP.  
 AC ABR6573;

DT 24-SEP-2002 (first entry)

DE cDNA encoding human ether-a-go-go related interacting protein MiRP1.

XX Human; human ether-a-go-go related gene; HERG; KCN1; MiRP1;  
 KW long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;  
 KW potassium channel; ss; gene.  
 XX  
 XX  
 OS Homo sapiens.  
 OS  
 XX

Key Location/Qualifiers  
 FT CDS 74..445  
 FT /\*tag= a  
 FT /product= "MiRP1"

PN W0200242735-A2.

PD 30-MAY-2002.

PF 30-OCT-2001; 2001MO-US45644.

PR 30-OCT-2000; 2000US-244340P.  
 XX  
 PA (UYVA-) UNIV VANDERBILT.  
 XX  
 PI Balser JR, George AL\*, Roden DM;  
 XX  
 DR WPI; 2002-527650/56.  
 XX P-PSDB: AAU99168.  
 PT Identifying a potassium channel activity modulator for drug design,  
 PT comprises contacting a compound with a potassium channel and rat  
 PT cerebellar cDNA library (KCRI) polypeptide, and determining activity -  
 XX  
 XX  
 PS Claim 17; Page 162-163; 164pp; English.  
 CC The invention relates to identifying (M1) a compound that modulates  
 CC biological activity of a potassium channel (PC), by contacting a  
 CC compound with a structure comprising a PC polypeptide and a polypeptide  
 CC cloned from a rat cerebellar cDNA library (KCRI), and determining the  
 CC activity of the PC polypeptide in the presence and absence of the  
 CC compound, where a difference in the activities indicates modulation of  
 CC biological activity of PC. Also include are identifying (M2) a candidate  
 CC compound that modulates the biological activity of a complex comprising a  
 CC human ether-a-go-go-related gene (HERG) channel polypeptide and a KCRI  
 CC polypeptide, identifying (M3) a candidate compound as a modulator of KCRI  
 CC expression, modulating (M4) PC function in a subject, comprising  
 CC administering to the subject a substance that provides expression of a  
 CC KCRI-encoding nucleic acid molecule in a cell or tissue, where modulated  
 CC PC function is desired, screening (M5) for susceptibility to a drug-  
 CC induced cardiac arrhythmia in a subject, comprising obtaining a  
 CC biological sample from the subject and detecting a polymorphism of a KCRI  
 CC gene in the biological sample from the subject, where the presence of the  
 CC polymorphism indicates the susceptibility of the subject to a  
 CC drug-induced cardiac arrhythmia, an oligonucleotide pair, where a first  
 CC oligonucleotide of the pair hybridises to a first portion of a KCRI gene  
 CC which includes a polymorphism of the KCRI gene, and the second  
 CC oligonucleotide of the pair hybridises to a second portion of the KCRI  
 CC gene that is adjacent to the first portion and a set of antisense  
 CC oligonucleotide primers, suitable for amplifying a portion of a KCRI gene  
 CC which includes a polymorphism of the KCRI gene, (M1) is useful for  
 CC identifying a compound that modulates biological activity of PC,  
 CC especially HERG, for modulating PC function (i.e. modulating HERG  
 CC activity) in a mammal, by preparing a composition comprising the  
 CC compound and administering the composition. The compound is useful for  
 CC treating or preventing long QT syndrome (LQT) and is useful in drug  
 CC designing. The present sequence encodes a HERG interacting  
 CC protein MiRP1 (not defined).  
 CC  
 XX  
 SQ Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match 99.8%; Score 730.4; DB 24; Length 732;  
 Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
 Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTGTCGGCTATTATTAATTAATGCA 60  
 DB 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTGTCGGCTATTATTAATTAATGCA 60  
 QY 61 GCAGAGGGAGAGCATGCTACTTATTCATTTGACACAGAGCGTGAAGAGCTTCCG 120  
 DB 61 GCAGAGGGAGAGCATGCTACTTATTCATTTGACACAGAGCGTGAAGAGCTTCCG 120  
 QY 121 AAGGATTTTATCTATATGACAAATGGGCGCAAGAACAAAGCTGACAGAGGC 180  
 DB 121 AAGGATTTTATCTATATGACAAATGGGCGCAAGAACAAAGCTGACAGAGGC 180  
 QY 181 CCTCCAGCCAAAGTGTGAGTAAGTCTGATGCTGATGCTGATGATGATGAT 240  
 DB 181 CCTCCAGCCAAAGTGTGAGTAAGTCTGATGCTGATGCTGATGATGATGATGAT 240  
 QY 241 GATTGGAATGTTCTTTCATCATCGTGCCATCCTGGTGAAGCACTGTGAATCCAGAG 300  
 DB 241 GATTGGAATGTTCTTTCATCATCGTGCCATCCTGGTGAAGCACTGTGAATCCAGAG 300

QY 301 ACGGGAACATCCATGACCCCTACACAGTACATGTAGAGGACGGCAGAAAAGTA 360  
 DB 301 ACGGGAACATCCATGACCCCTACACAGTACATGTAGAGGACGGCAGAAAAGTA 360  
 QY 361 CAAGAGCCAAATCTTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAACATTTGGTGC 420  
 DB 361 CAAGAGCCAAATCTTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAACATTTGGTGC 420  
 QY 421 GCGTGGGTTCAAAATGTCCTCCCTGATTAAGGAGAAAGCACCAGCAAGCTAGACGTC 480  
 DB 421 GCGTGGGTTCAAAATGTCCTCCCTGATTAAGGAGAAAGCACCAGCAAGCTAGACGTC 480  
 QY 481 CAGACATGAAGAGATGCCAGTGCACGAGGCAATTCAAATTTGCTTGTGATTAAGAAA 540  
 DB 481 CAGACATGAAGAGATGCCAGTGCACGAGGCAATTCAAATTTGCTTGTGATTAAGAAA 540  
 QY 541 GTGAGTTCCTTGTCTTGTGAGAAATTTTCATGAGATATGTGTGGCCATAAAGA 600  
 DB 541 GTGAGTTCCTTGTCTTGTGAGAAATTTTCATGAGATATGTGTGGCCATAAAGA 600  
 QY 601 TAGATGACATTTCAATCTCATGATTTATGCTGCTTGTGGACCAATATTTTGTCTGA 660  
 DB 601 TAGATGACATTTCAATCTCATGATTTATGCTGCTTGTGGACCAATATTTTGTCTGA 660  
 QY 661 AGACCTCTTTACATTTCCGGGCAAGTGAATGTCATTTTAAATCAATATGATGAAT 720  
 DB 661 AGACCTCTTTACATTTCCGGGCAAGTGAATGTCATTTTAAATCAATATGATGAAT 720  
 QY 721 AAAGCCAAATTT 732  
 DB 721 AAAGCCAAATTT 732

RESULT 8  
 AAD35169  
 ID AAD35169 standard; DNA; 732 BP.  
 XX  
 AC AAD35169;  
 XX  
 DT 25-JUN-2002 (first entry)  
 XX  
 DE Human KCNE2 mutant DNA (C420T).  
 XX  
 KW Human; Min-K related ion channel protein; MiRP1; ion channel disorder;  
 KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;  
 KW single nucleotide polymorphism; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key location/Qualifiers  
 FT CDS 74..445  
 FT /\*tag= a /product= "Human MiRP1 mutant protein"  
 FT variation /replace (420, C)  
 FT /\*tag= b /standard\_name= "single nucleotide polymorphism (SNP)"  
 PN W0200222875-A2.  
 PD 21-MAR-2002.  
 PE 11-SEP-2001; 2001WO-US28332.  
 PR 11-SEP-2000; 2000US-231571P.  
 PA (UYVA ) UNIV YALE.  
 PI Goldstein SAN;  
 XX  
 XX WPI; 2002-362360/39.  
 DR P-PSDB: AAE22094.  
 XX

PT Novel gene encoding Min-K related ion channel protein subunit and  
PT polymorphisms in this gene associated with antibiotic-induced long QT  
syndrome -

PS Claim 1; Page 41-42; 49pp; English.

CC The present invention relates to novel KCNE2 genes encoding Min-K related  
CC (MiRP) 1 ion channel proteins and polymorphisms in these genes that are  
CC associated with ion channel disorders including antibiotic-induced long  
CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position  
CC encoding the amino acid positions is useful for diagnosing the presence  
CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
CC are useful in the development of new drug therapies which selectively  
CC target one or more KCNE2 polymorphisms that are associated with cardiac  
CC arrhythmias. The present sequence is human KCNE2 mutant DNA (C420T).

SO Sequence 732 BP; 221 A; 151 C; 157 G; 203 T; 0 other;

Query Match 99.8%; Score 730.4; DB 24; Length 732;  
Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCGCTATTATTTAAATTGCA 60  
DB 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCGCTATTATTTAAATTGCA 60  
QY 61 GCAGGAGGGAAGCATGCTACTTATTCAAATTCACACAGAGCTGGAAGACGCTCCG 120  
DB 61 GCAGGAGGGAAGCATGCTACTTATTCAAATTCACACAGAGCTGGAAGACGCTCCG 120  
QY 121 AAGGATTTTATTTACTTATATGACAAATTTGGCCGACAGACAGCTGAGAGGC 180  
DB 121 AAGGATTTTATTTACTTATATGACAAATTTGGCCGACAGACAGCTGAGAGGC 180  
QY 181 CCTCCAGGCCAAAGTGTATGCTGAGAACTTCTACTATGCTGATCCATGATGAT 240  
DB 181 CCTCCAGGCCAAAGTGTATGCTGAGAACTTCTACTATGCTGATCCATGATGAT 240  
QY 241 GATTGGAATGTTCTTTTCATCATCGTGGCCATCCTGCTGAGACACTGGAATCCAAAG 300  
DB 241 GATTGGAATGTTCTTTTCATCATCGTGGCCATCCTGCTGAGACACTGGAATCCAAAG 300  
QY 301 ACGGGAACACTCCCAATGACCCCTACCCACAGTACATTTGAGAGACTGCGAGAAAAGTA 360  
DB 301 ACGGGAACACTCCCAATGACCCCTACCCACAGTACATTTGAGAGACTGCGAGAAAAGTA 360  
QY 361 CAAGAGCCAAATCTTGAATCTAGAAAGATGGAAGGCCACCATCATGAGAACTGGTGC 420  
DB 361 CAAGAGCCAAATCTTGAATCTAGAAAGATGGAAGGCCACCATCATGAGAACTGGTGC 420  
QY 421 GCGTGGGTTCAAAATGTCCCTGATTAAGGAGAGAAAGCAACAAGCTAGACGTGACGTC 480  
DB 421 GCGTGGGTTCAAAATGTCCCTGATTAAGGAGAGAAAGCAACAAGCTAGACGTGACGTC 480  
QY 481 CAGACATGAAGAAGATGCCAGTGCACAGAGCAAAATCCAAATTTGCTTGTAGAGAAA 540  
DB 481 CAGACATGAAGAAGATGCCAGTGCACAGAGCAAAATCCAAATTTGCTTGTAGAGAAA 540  
QY 541 GTGAGTCCCTGGCTTGTGGAATTTTCATGAGAGATATGCTTGGCCCAATAAGA 600  
DB 541 GTGAGTCCCTGGCTTGTGGAATTTTCATGAGAGATATGCTTGGCCCAATAAGA 600  
QY 601 TAGATGACATTTCAATCTCAGTATTTATGCTTGTGAGCAATATTTTGTGCTGA 660  
DB 601 TAGATGACATTTCAATCTCAGTATTTATGCTTGTGAGCAATATTTTGTGCTGA 660  
QY 661 AGACCTCTTACTTCCGGGCAAGTGAATGCAATTTTAAATCAATCAATGAGAAAT 720  
DB 661 AGACCTCTTACTTCCGGGCAAGTGAATGCAATTTTAAATCAATCAATGAGAAAT 720  
QY 721 AAAGCCAAATTT 732  
DB 721 AAAGCCAAATTT 732

DB 721 AAAGCCAAATTT 732

RESULT 9

ID AAD35171 standard; DNA; 732 BP.

AC AAD35171;

DT 25-JUL-2002 (first entry)

XX Human KCNE2 mutant DNA (T234C).

XX Human; Min-K related ion channel protein; MiRP1; ion channel disorder;

KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;

XX single nucleotide polymorphism; ds.

OS Homo sapiens.

EH Key Location/Qualifiers

FT CDS 74..445

FT FT /tag= a

FT variation /product= "Human MiRP1 mutant protein"

FT FT /replace (234, T)

FT FT /tag= b

XX /standard\_name= "Single nucleotide polymorphism (SNP)"

XX MO200222875-A2.

XX 21-MAR-2002.

XX 11-SEP-2001; 2001WO-US28332.

XX 11-SEP-2000; 2000US-231571P.

XX (UYVA ) UNIV YALE.

XX Goldstein SAN;

XX WPI; 2002-362360/39.

XX P-PSDB; AAE22096.

XX Novel gene encoding Min-K related ion channel protein subunit and

PT polymorphisms in this gene associated with antibiotic-induced long QT

PT syndrome -

XX Claim 12; Page 44-45; 49pp; English.

XX The present invention relates to novel KCNE2 genes encoding Min-K related

CC (MiRP) 1 ion channel proteins and polymorphisms in these genes that are

CC associated with ion channel disorders including antibiotic-induced long

CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,

CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position

CC encoding the amino acid positions is useful for diagnosing the presence

CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods

CC are useful in the development of new drug therapies which selectively

CC target one or more KCNE2 polymorphisms that are associated with cardiac

CC arrhythmias. The present sequence is human KCNE2 mutant DNA (T234C).

SO Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match 99.8%; Score 730.4; DB 24; Length 732;  
Best Local Similarity 99.9%; Pred. No. 9.2e-203;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCGCTATTATTTAAATTGCA 60  
DB 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTTGTCGCTATTATTTAAATTGCA 60  
QY 61 GCAGGAGGGAAGCATGCTACTTATTCAAATTCACACAGAGCTGGAAGACGCTCCG 120  
DB 61 GCAGGAGGGAAGCATGCTACTTATTCAAATTCACACAGAGCTGGAAGACGCTCCG 120







XX	bone disorder; osteoporosis; vascular growth disorder;
KW	tissue regeneration; wound healing; infection; immune disorder;
KW	cell culture; drug screening; gene therapy; antiinflammatory;
KW	antistatic; antiarthritic; haemostatic; antiarteriosclerotic;
KW	cytostatic; osteopathic; vasotrophic; cardiac; antiviral; antibacterial;
XX	antifungal; vulnery; antitumor; ss.
OS	Homo sapiens.
PN	WO200157188-A2.
PD	09-AUG-2001.
XX	
PF	05-FEB-2001; 2001WO-US03800.
XX	
XX	03-FEB-2000; 2000US-0496914.
PR	27-APR-2000; 2000US-0560875.
XX	
PA	(HYSE-) HYSBO INC.
XX	
PI	Tiang YT, Liu C, Drmanac RT;
XX	
XX	WPI; 2001-457740/49.
DR	P-PSDB; ABB11948.
XX	
PT	Human proteins and DNA encoding sequences useful for preventing,
PT	treating or ameliorating a medical condition in a mammalian subject
PT	e.g., arthritis and cancer -
XX	
PS	Claim 1; Page 826; 1963pp; English.
XX	
CC	Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and
CC	sequences ABA08225-ABA09574 represent nucleic acids encoding them. The
CC	invention also relates to vectors and recombinant host cells comprising a
CC	nucleotide of the invention, methods of producing the novel polypeptides,
CC	antibodies against the polypeptides, methods of detecting the nucleotides
CC	or polypeptides in a sample, and methods of identifying compounds which
CC	bind to polypeptides of the invention. Although novel, many of the
CC	polypeptides of the invention have homology to known proteins, thereby
CC	giving an insight into their probable biological activities, and hence
CC	potential therapeutic applications. The polypeptides of the invention may
CC	have various activities, including cytokine, cell proliferation or cell
CC	differentiation activities; stem cell growth factor activity;
CC	hematopoiesis regulatory activity; tissue growth activity;
CC	immunomodulatory activity; activin- or inhibin-related activities;
CC	chemotactic or chemokinetic activities; haemostatic, thrombotic or
CC	thrombolytic activities; receptor or ligand activities; or may be
CC	involved in oncogenesis, cancer cell proliferation or metastasis.
CC	Depending on their biological activities, polypeptides and nucleotides of
CC	the invention are useful for preventing, treating or ameliorating medical
CC	conditions, e.g., by protein or gene therapy. Such conditions include
CC	cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell
CC	disorders), chronic inflammatory conditions (e.g., asthma or arthritis),
CC	proliferative retinopathy, atherosclerosis, coronary heart disease,
CC	arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal
CC	vascular growth. Polypeptides involved with tissue regeneration and
CC	repair (or nucleic acids encoding them) may be used to promote wound
CC	healing (e.g., of burns, incisions and ulcers), while those with
CC	immunomodulatory activities may be used in the treatment of viral,
CC	bacterial and fungal infections in addition to immune disorders.
CC	Polypeptides with growth factor activity may be used in cell cultures to
CC	promote cell growth. For example, such polypeptides may be used to
CC	manipulate stem cells in culture to give rise to neuroepithelial cells
CC	that can be used to augment or replace cells damaged by illness,
CC	autoimmune disease or accidental damage. The polypeptides and nucleotides
CC	may also be used in the diagnosis of the above conditions, and in drug
CC	screening techniques. The present sequence represents a cDNA encoding a
XX	novel human polypeptide of the invention.
XX	
SD	Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;
Query Match	75.7%; Score 553.8; DB 22; Length 600;
Best Local Similarity	99.6%; Pied NO. 2.7e-151;

	Matches	555; Conservative	0;	Mismatches	2;	Indels	0;	Gaps						
Qy	63	AGAGGAGAGCATTGCTACTTACTTATTCCAATTTACACAGACGCTGGAGAGCTTCCGAA	122											
Db	27	ATGAGGAGACATGCTGACTTTATTCCAATTTACACAGACGCTGGAGAGCTTCCGAA	86											
Qy	123	GGATTTTATTTACTTATTAATGACAAATTTGGCGGACAAACAAACGCTGAGCAAGAGGCC	182											
Db	87	GGATTTTATTTACTTATTAATGACAAATTTGGCGGACAAACAAACGCTGAGCAAGAGGCC	146											
Qy	183	TCCAAAGCCAAAGTGTGATGCTGAGAACCTTCTACTATGTCATCTGACTCTATGCTGATGA	242											
Db	147	TCCAAGCCAAAGTGTGATGCTGAGAACCTTCTACTATGTCATCTGACTCTATGCTGATGA	206											
Qy	243	TTTGAATGTTCTCTTTTCATCATCGTGGCCATCTCTGCTGAGCAGCTGTGAATCCAAAGAC	302											
Db	207	TTTGAATGTTCTCTTTTCATCATCGTGGCCATCTCTGCTGAGCAGCTGTGAATCCAAAGAC	266											
Qy	303	GGGAAACCTCCATGACCCCTACCCACATACATTTGTAGAGACCTGGAGAGAAAGTACA	362											
Db	267	GGGAAACCTCCATGACCCCTACCCACATACATTTGTAGAGACCTGGAGAGAAAGTACA	326											
Qy	363	AGAGCCAAATCTTGAATCTGAAAGATCGAAGGCCACCATCCATGAGAAATTTGGTCGG	422											
Db	327	AGAGCCAAATCTTGAATCTGAAAGATCGAAGGCCACCATCCATGAGAAATTTGGTCGG	386											
Qy	423	CTGGGTTCCAAATATGTCCTCCCTGATTAAGGAGAGAACCAAGCTAACATCTGACGTCA	482											
Db	387	CTGGGTTCCAAATATGTCCTCCCTGATTAAGGAGAGAACCAAGCTAACATCTGACGTCA	446											
Qy	483	GACATGAAGAGATGCCAGTCCACAGGCGCAAAATTCCTTCTTGTGTGAAGAAAGT	542											
Db	447	GACATGAAGAGATGCCAGTCCACAGGCGCAAAATTCCTTCTTGTGTGAAGAAAGT	506											
Qy	543	GAGTTCCTGCTCTTGTGAGAAATTTTCATGAGATTATGTTGGTCCAAATTAAGATA	602											
Db	507	GAGTTCCTGCTCTTGTGAGAAATTTTCATGAGATTATGTTGGTCCAAATTAAGATA	566											
Qy	603	GATGACATTTCAATCTC	619											
Db	567	GATGACATTTCAATCTC	583											
RESULT 13														
AAK52645														
ID	AAK52645	standard; cDNA; 600 BP.												
XX	AAK52645;													
AC														
XX	06-NOV-2001 (first entry)													
DT														
XX														
DE	Human polynucleotide SEQ ID NO 2174.													
XX														
KM	Human; cytokine; cell proliferation; cell differentiation; gene therapy;													
KW	vaccine; peptide therapy; stem cell growth factor; leukemia;													
KW	tissue growth factor; immunomodulatory; cancer; leukemia;													
KW	nervous system disorder; arthritis; inflammation; ss.													
OS	Homo sapiens.													
XX														
PN	WO200157190-A2.													
XX														
PD	09-AUG-2001.													
XX														
PF	05-FEB-2001; 2001MO-US04098.													
XX														
PR	03-FEB-2000; 2000US-0496914.													
PR	27-APR-2000; 2000US-0560875.													
PR	20-JUN-2000; 2000US-0598075.													
PR	19-JUL-2000; 2000US-0620325.													
PR	01-SEP-2000; 2000US-0654936.													

PR 30-NOV-2000; 2000US-0728422.  
XX  
PA (HYSE-) HYSEQ INC.  
XX  
PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;  
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;  
XX  
DR WPI: 2001-476283/51.  
DR P-PSDB; AAM79512.  
XX  
PT Nucleic acids encoding polypeptides with cytokine-like activities,  
PT useful in diagnosis and gene therapy -  
XX  
PS Claim 1; Page 4539-4540; 6221pp; English.  
XX  
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX  
XX Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;  
XX  
Query Match 75.7%; Score 553.8; DB 22; Length 600;  
Best Local Similarity 99.6%; Pred. No. 2.7e-151;  
Matches 555; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
XX  
QY 63 AGAGGAGAGCATGTCCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAA 122  
DB 27 ATGAGGAGACATGTCCTACTTTATTCACAAATTCACACAGACGCTGGAAGAGCTTCCGAA 86  
QY 123 GGATTTTATTTACTTATATGAGACAATGGGGCCAGAACACAGCTGAGACAGAGGCC 182  
DB 87 GGATTTTATTTACTTATATGAGACAATGGGGCCAGAACACAGCTGAGACAGAGGCC 146  
QY 183 TCCAGCCAAAGTGTATGAGAACTTCTACTATGTATGTATCTGTACCTCATGGTATGA 242  
DB 147 TCCAGCCAAAGTGTATGAGAACTTCTACTATGTATGTATCTGTACCTCATGGTATGA 206  
QY 243 TTGGAAATGTTCTCTTTATCATGTGCGCATCTCTGTGAGCACTGTGAATCCAAAGAGAC 302  
DB 207 TTGGAAATGTTCTCTTTATCATGTGCGCATCTCTGTGAGCACTGTGAATCCAAAGAGAC 266  
QY 303 GGGAAACCTCCAAATGAGACCCCTACACACAGTACATTTGAGAGAGTGGCAGAGAAAGTACA 362  
DB 267 GGGAAACCTCCAAATGAGACCCCTACACACAGTACATTTGAGAGAGTGGCAGAGAAAGTACA 326  
QY 363 AGAGCCAAATCTTGAATCTGAGAAATCGAAGCCACATCCATCGAAGACATTTGGTGGC 422  
DB 327 AGAGCCAAATCTTGAATCTGAGAAATCGAAGCCACATCCATCGAAGACATTTGGTGGC 386  
QY 423 CTGGGTTCAAAATGTCCCTCTGTAAGGGAGAAAGCCAGCTAAACATCTGACGTCCA 482  
DB 387 CTGGGTTCAAAATGTCCCTCTGTAAGGGAGAAAGCCAGCTAAACATCTGACGTCCA 446  
QY 483 GACATGAGAGATGCCAGTCCAGAGGAGCAATCCAAATGTTCTTGGCTTGAAGAAAGT 542  
DB 447 GACATGAGAGATGCCAGTCCAGAGGAGCAATCCAAATGTTCTTGGCTTGAAGAAAGT 506  
QY 543 GAGTCTCTGCTCTTGTGAGATTTTCATGAGATTTATGTGGTGGCCAAATGAAGATA 602  
DB 507 GAGTCTCTGCTCTGCTGTTGAGATTTTCATGAGATTTATGTGGTGGCCAAATGAAGATA 566

QY 603 GATGACATTTCAATCTC 619  
DB 567 GATGACATTTCAATCTC 583  
XX  
RESULT 14  
AAK51661  
ID AAK51661 standard; cDNA; 655 BP.  
XX  
XX AAK51661;  
XX  
XX 06-NOV-2001 (first entry)  
XX  
DE Human polynucleotide SEQ ID NO 206.  
XX  
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;  
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;  
KW tissue growth factor; immunomodulatory; cancer; leukaemia;  
KW nervous system disorder; arthritis; inflammation; ss.  
XX  
OS Homo sapiens.  
PN MO200157190-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 05-FEB-2001; 2001WO-US04098.  
XX  
PR 03-FEB-2000; 2000US-0496914.  
PR 27-APR-2000; 2000US-0560875.  
PR 20-JUN-2000; 2000US-0598075.  
PR 19-JUL-2000; 2000US-0620325.  
PR 01-SEP-2000; 2000US-0654936.  
PR 15-SEP-2000; 2000US-0663561.  
PR 20-OCT-2000; 2000US-0693325.  
PR 30-NOV-2000; 2000US-0728422.  
XX  
PA (HYSE-) HYSEQ INC.  
XX  
PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;  
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;  
XX  
DR WPI: 2001-476283/51.  
DR P-PSDB; AAM78528.  
XX  
PT Nucleic acids encoding polypeptides with cytokine-like activities,  
PT useful in diagnosis and gene therapy -  
XX  
PS Claim 1; Page 1024; 6221pp; English.  
XX  
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX  
XX Sequence 655 BP; 196 A; 154 C; 146 G; 153 T; 6 other;  
XX  
Query Match 75.7%; Score 553.8; DB 22; Length 655;  
Best Local Similarity 99.6%; Pred. No. 2.9e-151;  
Matches 555; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
QY 63 AGAGGAGAGCATGTCCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAA 122

```

Db      82 ATGAGGAGAGATGCTACTTATCCAAATTCACACAGAGCTGGAGAGCTCTCCGAA 141
QY      123 GGATTTTATTAATTAATGACAAATTTGGCCGAGAAACACAGACTGACCAAGAGCCC 182
Db      142 GGATTTTATTAATTAATGACAAATTTGGCCGAGAAACACAGACTGACCAAGAGCCC 201
QY      183 TCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGCTACCTATGCTATGA 242
Db      202 TCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGCTACCTATGCTATGA 261
QY      243 TTGGAATGTTCTCTTTTATCATGCTGGCCATCCGCTGAGACACTGTGAATTCGAAGAC 302
Db      262 TTGGAATGTTCTCTTTTATCATGCTGGCCATCCGCTGAGACACTGTGAATTCGAAGAC 321
QY      303 GGGAAACCTCCAAATGACCCCTACACCAAGTACATTTGAGAGAGCTGGCAGAAAAGTACA 362
Db      322 GGGAAACCTCCAAATGACCCCTACACCAAGTACATTTGAGAGAGCTGGCAGAAAAGTACA 381
QY      363 AGAGCCAAATCTGATCTGATGAAGAAATCGAAGGCCACCATCCATGAGAAACATTGTGCGG 422
Db      382 AGAGCCAAATCTGATCTGATGAAGAAATCGAAGGCCACCATCCATGAGAAACATTGTGCGG 441
QY      423 CTGGGTTCAAAATGTCCCTCGATGAAGGAGAAAGCCACCAAGCTACATCTGACGTCCA 482
Db      442 CTGGGTTCAAAATGTCCCTCGATGAAGGAGAAAGCCACCAAGCTACATCTGACGTCCA 501
QY      483 GACATGAAGAGATGCCAGTGGCCAGAGGCAAAATCCAAATTTGCTTTGCTTGAAGAAAGT 542
Db      502 GACATGAAGAGATGCCAGTGGCCAGAGGCAAAATCCAAATTTGCTTTGCTTGAAGAAAGT 561
QY      543 GAGTTCCTTGCTCTTTTGTGGAATTTTCATGGAGATTAATGTGGTGGCCAAATTAAGATA 602
Db      562 GAGTTCCTTGCTCTTTGGAATTTTCATGGAGATTAATGTGGTGGCCAAATTAAGATA 621
QY      603 GATGACATTTCAATCTC 619
Db      622 GATGACATTTCAATCTC 638

RESULT 15
AAF80269
ID      AAF80269 standard; DNA; 471 BP.
XX
AC      AAF80269;
XX
DT      29-JUN-2001 (first entry)
XX
DE      Nucleotide sequence of human potassium channel subunit Isk2.
XX
KW      Human; potassium channel; Isk2; gene therapy; gastric motility;
KW      gastric acid secretion; anti-arrhythmic agent; myocardial infarction; ss.
XX
OS      Homo sapiens.
XX
XX
FH      Key      Location/Qualifiers
FT      CDS      79..450
FT      /tag= a
FT      /product= "potassium channel subunit Isk2"
XX
PN      WO200127246-A1.
XX
PD      19-APR-2001.
XX
PF      10-OCT-2000; 2000WO-US28014.
XX
PR      12-OCT-1999; 99US-0158781.
XX
PA      (MERI ) MERCK & CO INC.
XX
PI      Swanson RJ, Liu Y, Folander K;
XX
DR      WPI; 2001-273764/28.

```

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DR      P-PSDB; AAB67800.
XX
XX      New DNA encoding the Isk2 potassium channel subunit, useful e.g. for
PT      detecting mutations and screening for therapeutic agents
XX
XX      Claim 3; Fig 1A; 46pp; English.
XX
CC      The present sequence encodes a human potassium channel subunit,
CC      designated Isk2. The Isk2 polynucleotide, and derived probes, are
CC      used diagnostically to detect mutations in the Isk2 gene, to determine
CC      levels of mRNA expression and to isolate homologous sequences; for
CC      recombinant expression of Isk2; in gene therapy to increase potassium
CC      channel activity and to generate transgenic animals, as models and
CC      for drug screening. Recombinant Isk2 is used for studying biochemical
CC      activity of Isk2 and its role in disorders of gastric motility and
CC      gastric acid secretion, and to raise specific antibodies. Isk2
CC      modulators are potentially useful for treating diseases associated with
CC      increased or reduced potassium channel activity, e.g. as
CC      anti-arrhythmic agents for treating myocardial infarction and as
CC      regulators of gastric acid secretion.
XX
SQ      Sequence 471 BP; 143 A; 110 C; 103 G; 115 T; 0 other;

Query Match      63.7%; Score 466; DB 22; Length 471;
Best Local Similarity 100.0%; Pred. No. 1e-125;
Matches 466; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CAAATCCAGAAAAGATCCGTTTCTTACCTTGTCGCTATTTATTAATTAATGA 60
Db      6 CAAATCCAGAAAAGATCCGTTTCTTACCTTGTCGCTATTTATTAATTAATGA 65
QY      61 GCGAGGAGGAGATGTCATCTTATCCAAATTCACACAGAGCTGGAGAGCTCTCCG 120
Db      66 GCGAGGAGGAGATGTCATCTTATCCAAATTCACACAGAGCTGGAGAGCTCTCCG 125
QY      121 AAGGATTTTATTAATTAATGAGCAATTTGGCCGAGAAACACAAAGCTGAGCAAGAGC 180
Db      126 AAGGATTTTATTAATTAATGAGCAATTTGGCCGAGAAACACAAAGCTGAGCAAGAGC 185
QY      181 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGCTACCTATGCTAT 240
Db      186 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCTCTGCTACCTATGCTAT 245
QY      241 GATTGAATGTTCTCTTTTATCATGCTGGCCATCCGCTGAGACCTGTAATCCAAAGAG 300
Db      246 GATTGAATGTTCTCTTTTATCATGCTGGCCATCCGCTGAGACCTGTAATCCAAAGAG 305
QY      301 ACGGAAACCTCCAAATGACCCCTACACCAAGTACATTTGAGAGACTGGCAGAAAAGTA 360
Db      306 ACGGAAACCTCCAAATGACCCCTACACCAAGTACATTTGAGAGACTGGCAGAAAAGTA 365
QY      361 CAAGAGCCAAATCTGATCTGAGAAATCGAAGAGCCACCATCCATGAGAAACATTTGCTGC 420
Db      366 CAAGAGCCAAATCTGATCTGAGAAATCGAAGAGCCACCATCCATGAGAAACATTTGCTGC 425
QY      421 GCGTGGGTTCAAAATGTCCCTGATTAAGGAGGAGCAACCAAGC 466
Db      426 GCGTGGGTTCAAAATGTCCCTGATTAAGGAGGAGCAACCAAGC 471

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Search completed: May 21, 2003, 20:27:26  
Job time : 183.18 secs



GenCore version 5.1.4.p5\_4578  
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# OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:44:14 ; Search time 1593.65 Seconds

(without alignments)  
13367.622 Million cell updates/sec

Title: US-09-550-163-1

Perfect score: 732  
Sequence: 1 caaatccagaagaatccgt.....atgaataaagccaattc 732

Scoring table:

IDENTITY\_NUC  
Gapop 10.0 ; Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :  
1: GenEmbl:\*  
2: gb\_da:\*  
3: gb\_htg:\*  
4: gb\_in:\*  
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10: gb\_ro:\*  
11: gb\_sts:\*  
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13: gb\_un:\*  
14: gb\_vl:\*  
15: em\_da:\*  
16: em\_fun:\*  
17: em\_hum:\*  
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19: em\_mu:\*  
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21: em\_or:\*  
22: em\_ov:\*  
23: em\_pat:\*  
24: em\_ph:\*  
25: em\_pl:\*  
26: em\_ro:\*  
27: em\_sts:\*  
28: em\_un:\*  
29: em\_vl:\*  
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32: em\_htg\_inv:\*  
33: em\_htg\_mus:\*  
34: em\_htg\_pln:\*  
35: em\_htg\_rtd:\*  
36: em\_htg\_mam:\*  
37: em\_htg\_vrt:\*  
38: em\_sy:\*  
39: em\_htgo\_hum:\*  
40: em\_htgo\_mus:\*  
41: em\_htgo\_other:\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	732	100.0	732	6	AX406941 Sequence
2	732	100.0	732	9	AF071002 Homo sapi
3	730.4	99.8	732	6	AX406939 Sequence
4	730.4	99.8	732	6	AX406943 Sequence
5	730.4	99.8	732	6	AX406945 Sequence
6	730.4	99.8	732	6	AX406947 Sequence
7	730.4	99.8	24608	9	AP0000320 Homo sapi
8	730.4	99.8	100000	9	AP0000052 Homo sapi
9	730.4	99.8	100000	9	AP0000167 Homo sapi
10	730.4	99.8	100000	17	AP000120 Homo sapi
11	730.4	99.8	340000	9	AP001719 Homo sapi
12	671	91.7	809	9	AF302095 Homo sapi
13	292.2	39.9	1664	10	BC022659 Mus muscu
14	277.4	37.9	468	10	AF071003 Rattus no
15	273	37.3	144709	2	AC117904 Rattus no
16	266.4	36.4	372	10	AY050513 Cavia por
17	186.6	25.5	225	4	AY079211 Sus scrof
18	186.2	25.4	215	4	AF329636 Oryctolag
19	178.2	24.3	228	4	AF387764 Equus cab
20	56	7.7	534	4	RAB1PCS Oryctolag
21	55	7.5	750	10	GPIISK L20462 Cavia cobay
22	53.2	7.3	390	9	AF135188 Homo sapi
23	53.2	7.3	390	10	AY050512 Cavia por
24	53.2	7.3	398	6	I40373 Sequence 5
25	53.2	7.3	402	9	HDMISKA L33815 Homo sapien
26	53.2	7.3	408	9	HDMCDPCA L28168 Homo sapien
27	53.2	7.3	436	9	HUMISK M26685 Human ISK p
28	53.2	7.3	1703	6	AR119312 Sequence
29	53.2	7.3	3173	9	BC036452 Homo sapi
30	53.2	7.3	43126	9	AP0000324 Homo sapi
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32	53.2	7.3	100000	9	AP0000168 Homo sapi
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36	52.8	7.2	393	4	AB032575 Sus scrof
37	52.6	7.2	471	10	RATDRTRCA M36461 Rat delayed
38	51	7.0	585	10	RATPCPA M2412 Rat potassi
39	50.8	6.9	153306	2	AL807829 Danio rer
40	50.4	6.9	390	4	MSU62404 Felis catus
41	50.4	6.1	422	12	SYNMINK M64922 Synthetic v
42	44.6	5.9	209598	2	AC099473 Rattus no
43	43.2	5.9	209598	2	AC099473 Rattus no
44	43.2	5.8	122720	3	AC025721 Caenorhab
45	42.4	5.8	122720	3	AC025721 Caenorhab

## ALIGNMENTS

RESULT 1	AX406941	732 bp	DNA	linear	PAT 14-JUN-2002
LOCUS	AX406941	Sequence 3 from Patent WO0222875.			
DEFINITION	AX406941				
ACCESSION	AX406941.1	GI:21439816			
VERSION					
KEYWORDS					
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Goldstein, S.A.				
AUTHORS	Polymorphisms associated with cardiac arrhythmia				
TITLE	Patent: WO 0222875-A 3 21-MAR-2002;				
JOURNAL					

YALE UNIVERSITY (US)  
 Location/Qualifiers  
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 LEESKATIHENIGAGFKMSP"  
 BASE COUNT 221 a 152 c 157 g 202 t  
 ORIGIN

Query Match 100.0%; Score 732; DB 6; Length 732;  
 Best Local Similarity 100.0%; Pred. No. 4.6e-200;  
 Matches 732; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAATCCAGAAAAGATCCGTTTCCTAACCTGTGCGCTATTATTATTAAATTGCA 60  
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 QY 61 GCAGGAGGAGAGCATGCTCTACTTATTCGATTCACACAGACGCTGGAAGCTTCCG 120  
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 Db 121 AAGGATTTTATTACTATATGAGCAATTTGGCCGAGAACACAGAGCTGAGCAAGAGC 180  
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 Db 181 CCTCCAAAGCCAAAGTGTATGCTGAGAACTTCTACTATGTCATCCTGACTCATGTGAT 240  
 QY 241 GATTGGAATTTCTCTTTCATCATCGTGGCCATCCGTGGTGAAGACGTGAAATCCAAAG 300  
 Db 241 GATTGGAATTTCTCTTTCATCATCGTGGCCATCCGTGGTGAAGACGTGAAATCCAAAG 300  
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 Db 361 CAAGAGGCAATCTTGATCTAGAGAGATGAAAGGCCACATCCATGAGAACTTGGTGC 420  
 QY 421 GCGTGGGTTCAAAATGCCCCCTGATTAAGGAGAAAAGGCAAGGCTAACATCTGACGTC 480  
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 Db 721 AAAGCCAAATTT 732

RESULT 2  
 AF071002 732 bp mRNA linear PRI 29-APR-1999  
 LOCUS Homo sapiens mink-related peptide 1 mRNA, complete cds.  
 DEFINITION AF071002  
 ACCESSION AF071002.1 GI:4704422  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM  
 Homo sapiens.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 REFERENCE  
 1 (bases 1 to 732)  
 Abbott,G.W., Sesti,F., Splawski,I., Buck,M.E., Lehmann,M.H.,  
 Timothy,K.W., Keating,M.T. and Goldstein,S.A.  
 MIRP1 forms IKr potassium channels with HERG and is associated with  
 cardiac arrhythmia  
 Cell 97 (2), 175-187 (1999)  
 JOURNAL 99235979  
 MEDLINE 10219239  
 PUBMED 2 (bases 1 to 732)  
 REFERENCE  
 2 (bases 1 to 732)  
 Abbott,G.W., Sesti,F., Buck,M.E. and Goldstein,S.A.N.  
 Direct Submission  
 Submitted (05-JUN-1998) Section of Developmental Biology and  
 Biophysics, Department of Pediatrics and Boyer Center for Molecular  
 Medicine, Yale University School of Medicine, 295 Congress Avenue,  
 New Haven, CT 06536, USA  
 Location/Qualifiers

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 BASE COUNT 221 a 152 c 157 g 202 t  
 ORIGIN

Query Match 100.0%; Score 732; DB 9; Length 732;  
 Best Local Similarity 100.0%; Pred. No. 4.6e-200;  
 Matches 732; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAATCCAGAAAAGATCCGTTTCCTAACCTGTGCGCTATTATTATTAAATTGCA 60  
 Db 1 CAATCCAGAAAAGATCCGTTTCCTAACCTGTGCGCTATTATTATTAAATTGCA 60  
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 Db 61 GCAGGAGGAGAGCATGCTCTACTTATTCGATTCACACAGACGCTGGAAGCTTCCG 120  
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 Db 121 AAGGATTTTATTACTATATGAGCAATTTGGCCGAGAACACAGAGCTGAGCAAGAGC 180  
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QY 361 CAGAGCCAAATCTGAATCTAGAGATCGAAGGCCACCATTCATGAGACATTTGGTC 420  
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DB 601 TAGATGACATTTCAATCTCAGTGAATTAATGCTTGTGTTGAGCAATATTTTGTCTGA 660  
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DB 721 AAAGCCAAATTT 732

RESULT 3  
AX406939 732 bp DNA linear PAT 14-JUN-2002

LOCUS AX406939  
DEFINITION Sequence 1 from Patent WO0222875.  
ACCESSION AX406939  
VERSION AX406939.1 GI:21439814

KEYWORDS  
SOURCE

ORGANISM human.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1  
AUTHORS Goldstein, S.A.  
TITLE Polymorphisms associated with cardiac arrhythmia  
JOURNAL Patent: WO 0222875-A 1 21-MAR-2002;  
YALE UNIVERSITY (US)

FEATURES  
source location/Qualifiers  
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CDS  
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/replace="c"  
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BASE COUNT 221 a 151 c 157 g 203 t  
ORIGIN

Query Match 99.8%; Score 730.4; DB 6; Length 732;  
Best Local Similarity 99.9%; Pred. No. 1.3e-199;  
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 61 GCAGAGGAGAACGATGCTACTTATTCAAATTTACACAGACGCTGGAAGACGTTCTCG 120  
QY 121 AAGATTTTATTAATTAATATGACAAATTTGGCCGAGACACAAACGCTGAGCAAGGC 180  
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DB 121 AAGATTTTATTAATTAATATGACAAATTTGGCCGAGACACAAACGCTGAGCAAGGC 180  
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DB 181 CCTCAAGCCAAATGATGCTGAGACCTCTACTATGATCATCTGACTCATGGTAT 240  
QY 241 GATTGGAATGTTCTTTCATCATCTGAGCCATCTGTTGAGCACTGTGAAATCCAAGAG 300  
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DB 241 GATTGGAATGTTCTTTCATCATCTGAGCCATCTGTTGAGCACTGTGAAATCCAAGAG 300  
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DB 301 ACGGGAACACTCCATATACCCCTACCAACAGTACATTTGTAGAGACTGGCAGAGAAAGTA 360  
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DB 361 CAAAGGCCAAATCTTGAATCTAGAGAAATCGAAGGCCACCATTCATGAGACATTTGGTC 420  
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QY 481 CAGACATAGAGATGCCAGAGGCCACGAGCAATCCAAATTTGCTTGTGTTAAGAA 540  
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RESULT 4  
AX406943 732 bp DNA linear PAT 14-JUN-2002

LOCUS AX406943  
DEFINITION Sequence 5 from Patent WO0222875.  
ACCESSION AX406943  
VERSION AX406943.1 GI:21439818

KEYWORDS  
SOURCE

ORGANISM human.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1  
AUTHORS Goldstein, S.A.  
TITLE Polymorphisms associated with cardiac arrhythmia  
JOURNAL Patent: WO 0222875-A 5 21-MAR-2002;  
YALE UNIVERSITY (US)

FEATURES  
source location/Qualifiers  
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CDS  
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QY 661 AGACCTCTTTTACTTTCGCGGCAAGTGAATGTCATTTTAAATCAATCATGATGAAAT 720
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QY 721 AAGGCCAAATTT 732
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Db 721 AAGGCCAAATTT 732

RESULT 6
AX406947
LOCUS AX406947 732 bp DNA linear PAT 14-JUN-2002
DEFINITION Sequence 9 from Patent WO0222875.
ACCESSION AX406947
VERSION AX406947.1 GI:21439822
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
1 Goldstein, S.A.
AUTHORS Polymorphisms associated with cardiac arrhythmia
TITLE Patent: WO 0222875-A 9 21-MAR-2002;
JOURNAL YALE UNIVERSITY (US)
FEATURES
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location/Qualifiers
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variation
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Query Match 99.8%; Score 730.4; DB 6; Length 732;
Best Local Similarity 99.9%; Pred. No. 1.3e-199;
Matches 731; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTGTGCTTATTTATTTAAATTGCA 60
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Db 1 CAAATCCAGAAAAGATCCGTTTCTTAACCTGTGCTTATTTATTTAAATTGCA 60
QY 61 GCAGGAGGAGAGCATGCTACTTATTCATTCACACAGAGCCCTGGAGACCTCTCCG 120
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Db 61 GCAGGAGGAGAGCATGCTACTTATTCATTCACACAGAGCCCTGGAGACCTCTCTCG 120
QY 121 AAGGATTTTATTTACTTATATGACAAATTTGGCCAGAAACACAGCTGAGCAAGGC 180
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Db 121 AAGGATTTTATTTACTTATATGACAAATTTGGCCAGAAACACAGCTGAGCAAGGC 180
QY 181 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCCTGATCTGAT 240
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Db 181 CCTCCAAAGCCAAAGTTGATGCTGAGAACTTCTACTATGTCATCCCTGATCTGAT 240
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Db 241 GATTGGAATGCTCTTTGATCATCGTGGCCATCGCTGAGACAGCTGAAAACCAAGAG 300
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    |||
Db 361 CAAAGGCCAAATCTTGAATCTTAGAAGAAATCGAAGCCACCATTCATGAGAAATTTGTGC 420
QY 421 GGCTGGGTTCCAAATGTCGCCCTGATTAAGGGGAAAGGACACCAAGCTAAATCTGAGCTC 480
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Db 421 GGCTGGGTTCCAAATGTCGCCCTGATTAAGGGGAAAGGACACCAAGCTAAATCTGAGCTC 480
QY 481 CAGACATGAAAGATGCGACAGTCCACAGAGCAAAATCCAAATTTGCTTTAGAGAAA 540
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Db 661 AGACCTCTTTTACTTTCGCGGCAAGTGAATGTCATTTTAAATCAATCATGATGAAAT 720
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RESULT 7
AP000320
LOCUS AP000320 24608 bp DNA linear PRI 20-NOV-1999
DEFINITION Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AML region,
clone:Q12C8, complete sequence.
ACCESSION AP000320
VERSION AP000320.1 GI:4835689
KEYWORDS HTG.
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
REFERENCE
1 (bases 1 to 24608)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y. and Sakaki, Y.
Homo sapiens 24,608bp genomic DNA of 21q22.1
Published Only in Database (1999)
2 (bases 1 to 24608)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y. and Sakaki, Y.
Direct Submission
Submitted (13-MAY-1999) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gsr.riken.go.jp,
URL:http://ngs.gsc.riken.go.jp/, Tel:81-42-778-9923,
Fax:81-42-778-9924)
The sequence is a part of the data (ACCESSION No. AP000165 -
AP000173).
The sequencing project is supported by Japan Science Technology
Corporation (JST) and The Institute of Physical and Chemical
Research (RIKEN).
FEATURES
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 VERSION APO00052.1  
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 REFERENCE 1 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shibata,T. and Sakaki,Y.  
 TITLE Homo sapiens genomic DNA, chromosome 21q

JOURNAL Published Only in DataBase (1998)  
 REFERENCE 2 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shibata,T. and Sakaki,Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (11-MAY-1998) Masahira Hattori, Kitasato University,  
 Department of Science, JST Sequencing Laboratory; Kitasato 1-15-1,  
 Sagamihara 228, Japan (E-mail:hattori@hgc.ims.u-tokyo.ac.jp,  
 Tel.0427-78-9732, Fax:0427-78-9561)  
 COMMENT This sequence is conducted by Kitasato University JST sequencing  
 Laboratory as a JST sequencing team.  
 Principal Investigator:Toshiyuki Sakaki Ph.D.  
 Phone: +81-3-5449-5622, Fax: +81-3-5449-5445,  
 sakaki@hgc.ims.u-tokyo.ac.jp  
 Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D. The  
 sequence is submitted by Human Genome Sequencing in ALIS project of  
 JST

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clone B2344F14-f50E8, segment 3/9, complete sequence.
ACCESSION AP000167 GI:4827132
VERSION AP000167.1
KEYWORDS HTG.
SOURCE Homo sapiens DNA.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 100000)
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.
Homo sapiens 890,291bp genomic DNA of 21q22.1 (REGION: D21S226-AML
CLONE RANGE: B2344F14-f50E8)
2 (bases 1 to 100000)
Published only in Database (1999)
Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.
Direct Submission
Submitted (10-MAY-1999) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gsr.riken.go.jp,
URL:http://hsp.gsc.riken.go.jp/, Tel:81-42-778-9923,
Fax:81-42-778-9924)

COMMENT
E. coli transposon insertion: The present data does not contain E.
coli transposon sequences which integrated in the
original/previous sequences. We determined the boundary between
the insertion and genomic sequences experimentally, removed the
insertion sequences, reconstituted the present data. The sequencing
project is supported by Japan Science Technology Corporation (JST)
and The Institute of Physical and Chemical Research (RIKEN).

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DT 26-SEP-1999 (Rel. 61, Last updated, Version 3)
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DE region, segment 3/8.
XX
XX HTG.
XX
XX Homo sapiens (human)
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;
OC Eutheria; Primates; Catarrhini; Homiidae; Homo.
XX
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XX Hiraoka M., Yamaguchi H., Imai K., Shimada J.;
XX Submitted (15-Apr-1999) to the EMBL/Genbank/DDBJ databases.
XX Mita Hiraoka, Japan Science and Technology Corporation (JST), Advanced
XX Databases Department, 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081, Japan
XX (E-mail:mika@okyo.jst.go.jp, URL:http://www-alls.tokyo.jst.go.jp/,
XX Tel:81-3-5214-8491, Fax:81-3-5214-8470)

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RT  "Homo sapiens 817,199bp genomic DNA of 21q22.1 GART and AML region";
RL  Unpublished.
CC  This sequence is conducted by Kitasato University JST sequencing
CC  Laboratory as a JST sequencing team.
CC  Principal Investigator: Yoshiyuki Sakaki Ph.D.
CC  Phone: +81-3-5449-5622, Fax : +81-3-5449-5445,
CC  sakaki@qcc.ims.u-tokyo.ac.jp
CC  Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D.
CC  The sequence is submitted by Human Genome Sequencing in ALIS
CC  Project of JST.
CC  Japan Science and Technology Corporation (JST)
CC  5-3, Yonbancho, Chiyoda-Ku, Tokyo 102-0081 Japan
CC  For further information about this sequence, including its
CC  location and relationship to other sequences, please visit our
CC  sequence archive web site (http://www.alls.tokyo.jst.go.jp/MS/)
CC  or send email to webmaster@www.alls.tokyo.jst.go.jp
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Hattori M., Fujiyama A., Taylor T.D., Watanabe H., Yada T.,
Park H.S., Toyoda A., Ishii K., Totoki Y., Choi D.K., Soeda E.,
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Gardiner K., Nizetic D., Francis F., Lehrach H., Reinhardt R. and
Yaspo M.L.
The DNA sequence of human chromosome 21
Nature 405 (6784), 311-319 (2000)
20289799
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MEDLINE
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REFERENCE
AUTHORS

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TITLE  
 JOURNAL  
 Submitted (10-APR-2000) The Chromosome 21 Mapping and Sequencing Consortium: \* RIKEN Genomic Sciences Center, Human Genome Research Group \* Institute of Molecular Biotechnology, Genome Analysis \* Keio University School of Medicine, Dept. of Molecular Biology \* GBF, Dept. of Genome Analysis \* Max-Planck Institute for Molecular Genetics (addresses see below)



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KEYWORDS

SOURCE ORGANISM	REFERENCE AUTHORS TITLE	JOURNAL
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Homo sapiens.		
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1 (bases 1 to 809)		
Domenech,A., Estivill, X. and de la Luna,S.		
Cloning of human MIRP1 cDNA		
Unpublished		
2 (bases 1 to 809)		
Domenech,A., Estivill, X. and de la Luna,S.		
Direct Submission		
Submitted (01-SEP-2000) Medical and Molecular Genetics Center, Institut Recerca Oncologica, Avda. de Castelldefels Km 2,7', l'Hospitalet de Llobregat, Barcelona 08907, Spain		
Location/Qualifiers		

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ORIGIN	<p>           /protein_id="FACJ34.6.1"            /db_xref="GI:10121888"            /translation="MSTLSNFTOTLEDVFRRIPTITMDMWRONTTAEQELQAKVDAE            NEFVILVLMWIMGMSFTIIVALTIVTSVSKRREHNDPFIHOTIVEDWQEKYSQTLL            LEEKSTATHENIGAAAGKMPK"         </p>				

Query Match	91.7%	Score 671;	DB 9;	Length 809;
Best Local Similarity	99.7%	Pred. No. 1.9e-182;		
Matches 671; Conservative	1;	Mismatches	0;	Gaps 0;

Qy	60	AGCGSAGSGAAGCAATCTCTACTTTATCCAAATTTACACAGACAGCGTGAACACGCTTCC	119
Db	127	AGCAGSAGSGAAGCAATCTCTACTTTATCCAAATTTACACAGACAGCGTGAACACGCTTCC	186

0y	GAAGGATTTTAACTATTATGGAATTTGGCGCGAAGACCAACAGCTGAGGCAAGAG	179
120		
Db	GAAGGATTTTAACTATTATGGAATTTGGCGCGAAGACCAACAGCTGAGGCAAGAG	246
187		

OY	180	CCCTCCAGGCCAAAGTGTATGCTGTGAACACTTCTACTATGTGCATCCTGTACTTCATGGTGA	239
Dd	247	CCCTCCAGGCCAAAGTGTATGCTGTGAACACTTCTACTATGTGCATCCTGTACTTCATGGTGA	306

09	240	CGATTGGAAATCTTCTCTTTCATCATCGTGGCCATCCTGGAGACACTGTAAATCCAGA	299
Db	307	TGATTGGAATCTCTCTTTCATCATCGTGGCCATCCTGGAGACACTGTAAATCCAGA	366

QY	300	367	426
GACGGGAACACTCCATGACCCCTACACAGTACATTGTAGAGAGACTGGCAGGAAAGT	GACGGGAACACTCCATGACCCCTACACAGTACATTGTAGAGAGACTGGCAGGAAAGT	GACGGGAACACTCCATGACCCCTACACAGTACATTGTAGAGAGACTGGCAGGAAAGT	

QY 360 ACAAGAGCCAAATCTTGAATCTAGAGAATCGAAGCCACCATCATGAAACAATGGTG 419  
 427 ACAAGAGCCAAATCTTGAATCTAGAGAATCGAAGCCACCATCATGAGAACATTTGGTG 486  
 Db

QY	420	487	546
CGGCTGGGGTCAAAATGTCCCCCTATTAAGGAGAAAGCACAAGCTACATCTGAGCT	CGGCTGGGGTCAAAATGTCCCCCTATTAAGGAGAAAGCACAAGCTACATCTGAGCT	CGGCTGGGGTCAAAATGTCCCCCTATTAAGGAGAAAGCACAAGCTACATCTGAGCT	CGGCTGGGGTCAAAATGTCCCCCTATTAAGGAGAAAGCACAAGCTACATCTGAGCT

QY	480	547	606
CCAGCATGAGAGATGCCAGTGGCCACGAGGCAATCCAAATGCTTGGTTGAAGAA		CCAGCATGAGAGAGATGCCAGTGGCCACGAGGCAATCCAAATGCTTGGTTGAAGAA	

QY 540 AGTGGTTCCTTGCTTCCTTTGTGAAATATTCATGAGAGATTATGCGTGCCCAATAAG 599  
|||||  
DB 607 AGTGAATCCCTGCTCTCTGTGAAATATTCATGAGAGATTATGCGTGCCCAATAAG 666

[illegible]









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OM protein - protein search, using sw model

Run on: May 15, 2003, 14:15:58 ; Search time 13 Seconds

(without alignments)  
392.430 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632

Sequence: 1 MSTLSNFTQTEEDYFRRIFT.....ESKATIHENIGAGFKMSP 123

Scoring table:

BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_40:\*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	632	100.0	123	1	MIR1_HUMAN
2	554	87.7	123	1	MIR1_RAT
3	132.5	21.0	129	1	MIRK_HUMAN
4	130.5	20.6	129	1	MIRK_FELCA
5	128.5	20.3	129	1	MIRK_MOUSE
6	128	20.3	130	1	MIRK_MOUSE
7	127.5	20.2	130	1	MIRK_RAT
8	126.5	20.0	125	1	MIRK_PIG
9	125	19.8	130	1	MIRK_CAVPO
10	84.5	13.4	439	1	MIRK_RABIT
11	76.5	12.1	557	1	CPS_CLOTM
12	76	12.0	103	1	MIR2_HUMAN
13	74.5	11.8	1042	1	SYL_BORBU
14	74	11.7	938	1	VI20_HSVJ
15	73.5	11.6	497	1	NU2M_CHOCR
16	73	11.6	400	1	TYRP_HABIN
17	72	11.4	103	1	MIR2_MOUSE
18	71.5	11.3	107	1	MIR2_RAT
19	71	11.2	291	1	ATHB_RABIT
20	70.5	11.2	874	1	SVY_UREPA
21	70	11.1	170	1	MIR3_MOUSE
22	69.5	11.0	946	1	YBR6_YEAST
23	68.5	10.8	701	1	TP20_YEAST
24	68.5	10.8	1007	1	RGAL_YEAST
25	68	10.8	518	1	HEMK_RICPR
26	68	10.8	774	1	YMA6_YEAST
27	68	10.8	1175	1	SR82_YEAST
28	68	10.8	1607	1	MIRP_LYMS
29	67.5	10.7	453	1	YW5_CAEEL
30	67.5	10.7	1972	1	MYHB_RABIT
31	67	10.6	476	1	YG12_YEAST
32	66.5	10.5	752	1	PA26_MOUSE
33	66.5	10.5	1197	1	DPOH_PODAN

34	66	10.4	210	1	MT04_MYCGE
35	66	10.4	2146	1	INSR_DROME
36	65.5	10.4	987	1	EPB4_HUMAN
37	65.5	10.4	987	1	EPB4_MOUSE
38	65.5	10.4	2016	1	CIN5_HUMAN
39	65.5	10.4	2016	1	CIN5_HUMAN
40	65	10.3	2019	1	CIN5_RAT
41	65	10.3	622	1	VI57_CAEEL
42	65	10.3	629	1	ME25_SCHPO
43	65	10.3	1835	1	CCAI_RAT
44	64.5	10.2	556	1	YMC3_YEAST
45	64.5	10.2	809	1	PLSB_PASMU

## ALIGNMENTS

RESULT 1  
MIR1\_HUMAN STANDARD: PRT: 123 AA.  
AC Q916J6;  
DT 30-MAY-2000 (Rel. 39, Created)  
DT 30-MAY-2000 (Rel. 39, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE Minimum potassium ion channel-related peptide 1 (MIRP1) (Mink-related peptide 1).  
DE  
GN KCM2.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
OX NCBI\_TaxID=9606;  
[1]  
SEQUENCE FROM N.A., VARIANTS LQ76 E-9; T-54 AND T-57, AND VARIANT A-8.  
RC TISSUE-Heart;  
RX MEDLINE-99235979; PubMed-10219239;  
RA Abbott G.W., Seefelt F., Splawski I., Buck M.E., Lehmann M.H.,  
RA Timothy K.W., Keating M.T., Goldstein S.A.N.;  
RT "MIRP1 forms IKR potassium channels with HERG and is associated with cardiac arrhythmia.";  
RL Cell 97:175-187(1999).  
[2]  
SEQUENCE FROM N.A.  
RM Domenech A., Estivill X., de la Luna S.;  
RA "Cloning of human MIRP1 cDNA.";  
RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.  
[3]  
ASSOCIATION WITH KCMQ2/KCMQ3, AND TISSUE SPECIFICITY.  
RX MEDLINE-20487128; PubMed-11034315;  
RA Tinel N., Diocotot S., Lauritzen I., Barhanin J., Lazdunski M.,  
RA Borsotto M.;  
RT "M-type KCMQ2-KCMQ3 potassium channels are modulated by the KCM2 subunit.";  
RL FEBS Lett. 480:137-141(2000).  
-1- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE STABILITY OF THE MULTIMERIC COMPLEX. KCM2 CO-ASSEMBLE WITH KCMH2 (HERG) TO FORM THE CARDIAC POTASSIUM (IKR) CHANNEL. MAY ALSO ESSENTIALLY IDENTICAL PROPERTIES TO FORM A POTASSIUM CHANNEL WITH ASSOCIATE WITH KCMQ2 AND/OR KCMQ3 TO FORM A POTASSIUM CHANNEL WITH NATIVE M-CURRENT, A SLOWLY ACTIVATING AND DEACTIVATING POTASSIUM CONDUCTANCE WHICH PLAYS A CRITICAL ROLE IN DETERMINING THE SUBTHRESHOLD ELECTRICAL EXCITABILITY OF NEURONS. MAY ACT AS A REGULATORY SUBUNIT OF THE KCMQ2/KCMQ3 M-TYPE CHANNEL.  
-1- SUBUNIT: ASSOCIATES WITH KCMH2. MAY ASSOCIATE WITH KCMQ2 AND/OR KCMQ3.  
-1- SUBCELLULAR LOCATION: Type I membrane protein.  
-1- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN BRAIN, HEART, SKELETAL MUSCLE, PANCREAS, PLACENTA, KIDNEY, COLON AND THYMUS. A SMALL, BUT SIGNIFICANT EXPRESSION IS FOUND IN LIVER, OVARY, TESTIS, PROSTATE, SMALL INTESTINE AND LEUKOCYTES. VERY LOW EXPRESSION, NEARLY UNDETECTABLE, IN LONG AND SPLEEN.  
-1- DISEASE: DEFECTS IN KCM2 ARE THE CAUSE OF LONG QT SYNDROME TYPE 6 (LQ76); A FORM OF HEART DISEASE CHARACTERIZED BY CARDIAC

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CC ARRHYTHMIA. MUTANTS FORM CHANNELS THAT OPEN SLOWLY AND CLOSE
CC RAPIDLY, THEREBY DIMINISHING POTASSIUM CURRENTS.
CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
CC -1- DATABASE: NAME-IQTSdb; NOTE-KCNE2 mutations page;
CC WWW-http://www.ssi.dk/en/forakning/Iqtadb/kcne.htm".
CC -----
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CC -----
CC EMBL: AF071002; AAD28086.1; -.
CC EMBL: AF302095; AAG13416.1; -.
CC DR Genew; HGNC:6242; KCNE2.
CC DR MIM; 603796; -.
CC DR InterPro; IPR000369; ISK_Channel.
CC DR Pfam; PF02060; ISK_Channel; 1.
CC DR PRINTS; PR00168; KCNECHANNEL.
CC KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
CC Glycoprotein; Phosphorylation; Disease mutation; Polymorphism;
CC Long QT syndrome.
CC FT DOMAIN 1 48 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 49 69 POTENTIAL.
CC DOMAIN 70 123 CYTOPLASMIC (POTENTIAL).
CC CARBOHYD 6 6 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 29 29 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT MOD_RES 71 71 PHOSPHORYLATION (BY PKC) (POTENTIAL).
CC FT MOD_RES 74 74 PHOSPHORYLATION (BY PKC) (POTENTIAL).
CC FT VARIANT 8 8 T -> A.
CC FT VARIANT 9 9 /FTID-VAR_008375.
CC FT VARIANT 9 9 Q -> E (IN LQT6; IMPEDES ACTIVATION AND
CC INCREASES SENSITIVITY TO MACROLIDE
CC ANTIBIOTICS).
CC FT VARIANT 54 54 /FTID-VAR_008376.
CC FT VARIANT 54 54 M -> T (IN LQT6; FORMS I(KR) CHANNELS
CC THAT DEACTIVATE TWICE AS FAST AS WILD
CC TYPE).
CC FT VARIANT 57 57 /FTID-VAR_008377.
CC FT VARIANT 57 57 I -> T (IN LQT6).
CC FT VARIANT 57 57 /FTID-VAR_008378.
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Query Match 100.0%; Score 632; DB 1; Length 123;
Best Local Similarity 100.0%; Pred. No. 3.7e-56;
Matches 133; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MSTLSNFTQTLDEYFRIRITTYMDNMRONTAEQALQAKVDAENFYVILYIMWGMF 60
DB 1 MSTLSNFTQTLDEYFRIRITTYMDNMRONTAEQALQAKVDAENFYVILYIMWGMF 60
QY 61 SFTIIVALIVSVKSKRREHSNDPYHOYIYEDWQEKYSQILNLESKATIHENIGAAGFK 120
DB 61 SFTIIVALIVSVKSKRREHSNDPYHOYIYEDWQEKYSQILNLESKATIHENIGAAGFK 120
QY 121 MSP 123
DB 121 MSP 123
DB 121 MSP 123
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MIRL_RAT
ID MIRL_RAT STANDARD; PRT; 123 AA.
AC Q9MTW0;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DE 30-MAY-2000 (Rel. 39, Last annotation update)
DE Minimum potassium ion channel-related peptide 1 (MIRP1) (Mink-related
DE peptide 1).
GN KCNE2.
OS Rattus norvegicus (Rat).

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CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
CC NCBI_Taxid=10116;
CC [1]
CC SEQUENCE FROM N.A.
CC STRAIN-Sprague-Dawley; TISSUE=Heart;
CC MEDLINE=99235979; PubMed=10219239;
CC Abbott G.W., Sesti F., Splawski I., Buck M.E., Lehmann M.H.,
CC Timothy K.W., Keating M.T., Goldstein S.A.N.;
CC "MIRP1 forms IKR potassium channels with HERG and is associated with
CC cardiac arrhythmia."
CC Cell 97:175-187(1999)
CC -1- FUNCTION: ANCELLULAR PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM
CC CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE
CC STABILITY OF THE MULTIMERIC COMPLEX. KCNE2 CO-ASSEMBLE WITH KCNH2
CC (HERG) TO FORM THE CARDIAC POTASSIUM (IKR) CHANNEL.
CC -1- SUBUNIT: ASSOCIATES WITH KCNH2.
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF071003; AAD28087.1; -.
CC DR InterPro; IPR000369; ISK_Channel.
CC DR Pfam; PF02060; ISK_Channel; 1.
CC DR PRINTS; PR00168; KCNECHANNEL.
CC KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
CC Glycoprotein; Phosphorylation.
CC FT DOMAIN 1 48 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 49 69 POTENTIAL.
CC DOMAIN 70 123 CYTOPLASMIC (POTENTIAL).
CC CARBOHYD 6 6 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 29 29 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT MOD_RES 71 71 PHOSPHORYLATION (BY PKC) (POTENTIAL).
CC FT MOD_RES 74 74 PHOSPHORYLATION (BY PKC) (POTENTIAL).
CC SQ SEQUENCE 123 AA; 14356 MW; C91870E7B1EB82A CRC64;
Query Match 87.7%; Score 554; DB 1; Length 123;
Best Local Similarity 82.1%; Pred. No. 2.1e-48;
Matches 101; Conservative 17; Mismatches 5; Indels 0; Gaps 0;
QY 1 MSTLSNFTQTLDEYFRIRITTYMDNMRONTAEQALQAKVDAENFYVILYIMWGMF 60
DB 1 MSTLSNFTQTLDEYFRIRITTYMDNMRONTAEQALQAKVDAENFYVILYIMWGMF 60
QY 61 SFTIIVALIVSVKSKRREHSNDPYHOYIYEDWQEKYSQILNLESKATIHENIGAAGFK 120
DB 61 SFTIIVALIVSVKSKRREHSNDPYHOYIYEDWQEKYSQILNLESKATIHENIGAAGFK 120
QY 121 MSP 123
DB 121 MSP 123
DB 121 VSP 123
RESULT 3
MINK_HUMAN
ID MINK_HUMAN STANDARD; PRT; 129 AA.
AC P15382;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE 19k slow voltage-gated potassium channel protein (Minimal potassium
DE channel) (Mink).
GN KCNE1.
OS Homo sapiens (Human).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

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OK NCBI\_TaxID=9606;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RX MEDLINE-89273632; PubMed-2730656;  
 RA Murai T., Kakizuka A., Takumi T., Ohkubo H., Nakanishi S.;  
 RT "Molecular cloning and sequence analysis of human genomic DNA  
 RT encoding a novel membrane protein which exhibits a slowly activating  
 RT potassium channel activity";  
 RL Biochem. Biophys. Res. Commun. 161:176-181(1989).  
 RN [2]  
 RN SEQUENCE FROM N.A., AND VARIANT GLY-38.  
 RC TISSUE=Leukocyte;  
 RX MEDLINE-95129890; PubMed-7828904;  
 RA Lai L.P., Deng C.L., Moss A.J., Kass R.S., Liang C.S.;  
 RT "Polymorphism of the gene encoding a human minimal potassium ion  
 RT channel (minK).";  
 RL Gene 151:339-340(1994).  
 RN [3]  
 RN SEQUENCE FROM N.A.  
 RC TISSUE=Cornea;  
 RA Rae J.L.;  
 RT "Delayed rectifier potassium channel subunit from human cornea  
 RT epithelium";  
 RN Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.  
 RN [4]  
 RN VARIANT ASN-85.  
 RX MEDLINE-97055277; PubMed-8899564;  
 RA Tesson F., Douger C., Denjoy I., Berthet M., Benaacour M., Petit C.,  
 RA Coumel P., Schwartz K., Guicheney P.;  
 RT "Exclusion of KCNE1 (ISK) as a candidate gene for Jervell and  
 RT Lange-Nielsen syndrome.";  
 RL J. Mol. Cell. Cardiol. 28:2051-2055(1996).  
 RN [5]  
 RN TISSUE SPECIFICITY.  
 RX MEDLINE-97459933; PubMed-9312006;  
 RA Chouabe C., Neyroud N., Guicheney P., Lazdunski M., Romey G.,  
 RA Barhanin J.;  
 RT "Properties of Kv1Q1 K<sup>+</sup> channel mutations in Romano-Ward and Jervell  
 RT and Lange-Nielsen inherited cardiac arrhythmias.";  
 RL EMBO J. 16:5472-5479(1997).  
 RN [6]  
 RN VARIANT LQTS 58-PRO-PRO-59.  
 RX MEDLINE-97472471; PubMed-9328483;  
 RA Tyson J., Triebjerg L., Bellman S., Wren C., Taylor J.F.N.,  
 RA Batten J., Aslaksen B., Soerlund S.J., Lund O., Malcolm S.,  
 RA Pembrey M., Bhattacharya S., Blumer-Glindzic M.,  
 RT "ISK and Kv1Q1: mutation in either of the two subunits of the slow  
 RT component of the delayed rectifier potassium channel can cause  
 RT Jervell and Lange-Nielsen syndrome.";  
 RL Hum. Mol. Genet. 6:2179-2185(1997).  
 RN [7]  
 RN VARIANTS LQTS ILE-7 AND ASN-76.  
 RX MEDLINE-98016403; PubMed-9354783;  
 RA Schulze-Bahr E., Wang Q., Wedekind H., Haverkamp W., Chen O., Sun Y.,  
 RA Ruble C., Hordt M., Towbin J.A., Borggrete M., Assmann G., Qu X.,  
 RA Sonberg J.C., Breithardt G., Oberl C., Funke H.;  
 RT "KCNE1 mutations cause Jervell and Lange-Nielsen syndrome.";  
 RL Nat. Genet. 17:267-268(1997).  
 RN [8]  
 RN VARIANTS LQTS LEU-74 AND ASN-76.  
 RX MEDLINE-98016422; PubMed-9354802;  
 RA Splawski I., Tristani-Firouzi M., Lehmann M.H., Sanguinetti M.C.,  
 RA Keating M.T.;  
 RT "Mutations in the minK gene cause long QT syndrome and suppress ISK  
 RT function.";  
 RL Nat. Genet. 17:338-340(1997).  
 RN [9]  
 RN VARIANT LQTS ASN-76.  
 RX MEDLINE-98105943; PubMed-9445165;  
 RA Duggal P., Vesely M.R., Wattanasitichakoon D., Villafane J.,  
 RA Kaushik V., Beggs A.H.;  
 RT "Mutation of the gene for ISK associated with both Jervell and  
 RT Lange-Nielsen and Romano-Ward forms of Long-QT syndrome.";

RL Circulation 97:142-146(1998).  
 RN [10]  
 RN VARIANTS LQTS PHE-47, HIS-51, ASN-76 AND ARG-87.  
 RX MEDLINE-99330558; PubMed-10400998;  
 RA Bianchi L., Shen Z., Dennis A.T., Priori S.G., Napolitano C.,  
 RA Ronchetti E., Byskin R., Schwartz P.J., Brown A.M.;  
 RT "Cellular dysfunction of LQTS-minK mutants: abnormalities of ISKs, IKr  
 RT and trafficking in long QT syndrome.";  
 RL Hum. Mol. Genet. 8:1499-1507(1999).  
 RN [11]  
 RN VARIANTS LQTS HIS-32, TRP-98 AND THR-127.  
 RX MEDLINE-20432616; PubMed-10973849;  
 RA Splawski I., Shen J., Timothy K.W., Lehmann M.H., Priori S.G.,  
 RA Robinson J.L., Moss A.J., Schwartz P.J., Towbin J.A., Vincent G.M.,  
 RA Keating M.T.;  
 RT "Spectrum of mutations in long-QT syndrome genes. KV1Q1, HERG, SCN5A,  
 RT KCNE1, and KCNE2";  
 RL Circulation 102:1178-1185(2000).  
 RN [12]  
 RN VARIANT LQTS ILE-109.  
 RX MEDLINE-21546310; PubMed-11692163;  
 RA Schulze-Bahr E., Schwartz M., Haenschild S., Wedekind H., Funke H.,  
 RA Haverkamp W., Breithardt W., Pongs O., Isbrandt D., Breithardt G.;  
 RT "A novel long-QT 5 gene mutation in the C-terminus (V109I) is  
 RT associated with a mild phenotype.";  
 RL J. Mol. Med. 79:504-509(2001).  
 CC -I- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM  
 CC CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE  
 CC STABILITY OF THE MULTIMERIC COMPLEX. KCNE1 CO-ASSEMBLE WITH KCNQ1  
 CC (KV1Q1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC  
 CC POTASSIUM (IKs) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY  
 CC STATE ONLY AFTER 50 SECONDS.  
 CC -I- SUBUNIT: ASSOCIATES WITH KCNQ1.  
 CC -I- SUBCELLULAR LOCATION: type I membrane protein.  
 CC -I- TISSUE SPECIFICITY: EXPRESSED IN HEART, LUNG, KIDNEY, TESTIS,  
 CC OVARIES, SMALL INTESTINE, PERIPHERAL BLOOD LEUKOCYTES. NOT  
 CC DETECTED IN PANCREAS, SPLEEN, PROSTATE AND COLON. RESTRICTIVELY  
 CC LOCALIZED IN THE APICAL MEMBRANE PORTION OF EPITHELIAL CELLS.  
 CC -I- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY  
 CC SIMILARITY)  
 CC -I- DISEASE: DEFECTS IN KCNE1 ARE THE CAUSE OF THE AUTOSOMAL RECESSIVE  
 CC JERVELL AND LANGE-NIELSEN SYNDROME (JLNS). JLNS COMPRISES  
 CC PROFOUND CONGENITAL SENSORINEURAL DEAFNESS ASSOCIATED WITH  
 CC SYNCOPAL EPISODES. THESE ARE CAUSED BY VENTRICULAR TACHYARRHYTHMIA  
 CC SECONDARY TO ABNORMAL REPOLARIZATION, MANIFESTED BY A PROLONGED QT  
 CC INTERVAL ON THE ELECTROCARDIOGRAM.  
 CC -I- DISEASE: DEFECTS IN KCNE1 ARE THE CAUSE OF LONG QT SYNDROME TYPE 5  
 CC (LQTS5). A FORM OF HEART DISEASE CHARACTERIZED BY CARDIAC  
 CC ARRHYTHMIA. MUTANTS FORM CHANNELS THAT OPEN SLOWLY AND CLOSE  
 CC RAPIDLY, THEREBY DIMINISHING POTASSIUM CURRENTS.  
 CC -I- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.  
 CC -I- DATABASE: NAME=LQTSdb; NOTE=KCNE1 mutations page;  
 CC WWW="http://www.ssi.dk/en/forskning/lqtsdb/kcne1.htm".  
 CC -----  
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 CC -----  
 CC EMBL; M26685; AAA36129.1; -;  
 CC EMBL; L38815; AAA63905.1; -;  
 CC EMBL; L28168; AAA58418.1; -;  
 CC EMBL; AF135188; AAD25096.1; -;  
 CC PIR; A32447; A32447.  
 CC Genew; HGNC:6240; KCNE1.  
 CC MIM; 176261; -;  
 CC MIM; 220400; -;  
 CC InterPro; IPR000369; ISK\_Channel.  
 CC Pfam; PF02060; ISK\_Channel; 1.  
 CC PRINTS; PR00168; KCNECHANNEL.



```

RT mouse.*;
RL FEBS Lett. 301:168-172(1992).
CC -1- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM
CC CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE
CC STABILITY OF THE MULTIMERIC COMPLEX. KCNE1 CO-ASSEMBLE WITH KCNQ1
CC (KVLOT1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC
CC POTASSIUM (IKs) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY
CC STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).
CC -1- SUBUNIT: ASSOCIATES WITH KCNQ1 (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- TISSUE SPECIFICITY: RESTRICTIVELY LOCALIZED IN THE APICAL
CC MEMBRANE PORTION OF EPITHELIAL CELLS.
CC -1- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY
CC SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
CC -----
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CC -----
DR EMBL: X60457; CAA42990.1; -.
DR PIR: S17307; S17307.
DR PIR: S21135; S21135.
DR MGD: MGI:96673; Kcne1.
DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel.1.
DR PRINTS: PRO0168; KCNECHANNEL.
KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
KW Glycoprotein; Phosphorylation
FT DOMAIN 1 43 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 44 66 POTENTIAL.
FT DOMAIN 67 129 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 5 26 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT MOD_RES 102 102 PHOSPHORYLATION (BY PKC) (BY SIMILARITY).
SQ SEQUENCE 129 AA; 14578 MW; E66DF4742300E839 CRC64;

Query Match: 20.3%; Score 128.5; DB 1; Length 129;
Best Local Similarity 35.0%; Pred. No. 4.6e-06;
Matches 36; Conservative 22; Mismatches 28; Indels 17; Gaps 6;

QY 26 WQONTTAEQ-----EALQAKVDAENFYVILVMWIGKFSPTIIVAIIVSVKSKRR 77
DB 17 WCE--TAEGGVNSGLARKSQLRDQSK--LEALYILMWLGFEGFTLGLIMLSYRSKRL 71
QY 78 EHSNDPYHOYIYED-WQEKYS-QILNLESKAT-IHENIGA 116
DB 72 EHSHPENFYIESDAMQEKKAFFQARVLESFRACVIEHQAA 114

RESULT 6
MINK_RAT
ID MINK_RAT STANDARD; PRT; 130 AA.
AC P15383;
DT 01-APR-1990 (Rel. 14; Created)
DT 01-APR-1990 (Rel. 14; Last sequence update)
DT 16-OCT-2001 (Rel. 40; Last annotation update)
DE Isk slow voltage-gated potassium channel protein (Minimal potassium
DE channel) (Mink).
GN KCNE1.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Kidney;
RX MEDLINE=89058617; PubMed=3194754;
RA Takumi T., Ohkubo H., Nakanishi S.;

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RP "Cloning of a membrane protein that induces a slow voltage-gated
RT potassium current.*";
RL Science 242:1042-1045(1988).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Heart;
RX MEDLINE=90222152; PubMed=2183220;
RA Folander K., Smith J.S., Antanavage J., Bennett C., Stein R.B.,
RA Swanson R.;
RT "Cloning and expression of the delayed-rectifier Isk channel from
RT neonatal rat heart and diethylstilbestrol-primed rat uterus.*";
RL Proc. Natl. Acad. Sci. U.S.A. 87:2975-2979(1990).
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=uterus;
RX MEDLINE=90262731; PubMed=2344412;
RA Piragnell M., Snay R.J., Trimmer J.S., MacLusky N.J., Maffiolin F.,
RA Kaczmarek L.K., Boyle M.B.;
RT "Retrogen induction of a small, putative K+ channel mRNA in rat
RT uterus.*";
RL Neuron 4:807-812(1990).
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=91035347; PubMed=2229022;
RA Iwai M., Masu M., Tsuchida K., Mori T., Ohkubo H., Nakanishi S.;
RT "Characterization of gene organization and generation of
RT heterogeneous mRNA species of rat Isk protein.*";
RL J. Biochem. 108:200-206(1990).
RN [5]
RP MUTAGENESIS OF SER RESIDUES.
RX MEDLINE=92205350; PubMed=1553557;
RA Busch A.E., Varnum M.D., North R.A., Adelman J.P.;
RT "An amino acid mutation in a potassium channel that prevents
RT inhibition by protein kinase C.*";
RL Science 255:1705-1707(1992).
RN [6]
RP MUTAGENESIS OF ASP-77.
RX MEDLINE=95329285; PubMed=7605639;
RA Wang K.-W., Goldstein S.A.N.;
RT "Subunit composition of mink potassium channels.*";
RL Neuron 14:1303-1309(1995).
CC -1- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM
CC CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE
CC STABILITY OF THE MULTIMERIC COMPLEX. KCNE1 CO-ASSEMBLE WITH KCNQ1
CC (KVLOT1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC
CC POTASSIUM (IKs) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY
CC STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).
CC -1- SUBUNIT: ASSOCIATES WITH KCNQ1 (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- TISSUE SPECIFICITY: RESTRICTIVELY LOCALIZED IN THE APICAL
CC MEMBRANE PORTION OF EPITHELIAL CELLS.
CC -1- INDUCTION: BY ESTROGEN.
CC -1- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT.
CC -1- MISCELLANEOUS: MUTAGENESIS EXPERIMENTS WERE CARRIED OUT BY
CC EXPRESSING IN XENOPUS OOCYTES MINK MUTANT EITHER INDIVIDUALLY
CC (HOMOMULTIMERS) OR IN COMBINATION WITH WILD-TYPE MINK
CC (HETEROMULTIMERS) IN A 1:1 RATIO.
CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
CC -----
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CC -----
DR EMBL: M22412; AAA41822.1; -.
DR EMBL: M36461; AAA41098.1; -.
DR EMBL: D10709; BAA01553.1; ALT_SEQ.
DR PIR: A35633; A35633.
DR PIR: JH0140; JH0140.
DR PIR: A33177; A33177.

```

DR InterPro: IPR000369; ISK\_Channel.  
 DR Pfam: PF02060; ISK\_Channel; 1.  
 DR PRINTS: PR00168; KCNECHANNEL.  
 KM Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;  
 KM Glycoprotein; Phosphorylation.  
 FT DOMAIN 1 44 EXTRACELLULAR (POTENTIAL).  
 FT TRANSSEM 45 67 POTENTIAL.  
 FT DOMAIN 68 130 CYTOPLASMIC (POTENTIAL).  
 FT CARBOHYD 26 5 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 26 26 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT MOD\_RES 103 103 PHOSPHORYLATION (BY PKC) (PROBABLE).  
 FT MOTAGN 77 77 D->N: NO CURRENT (HOMOMULTIMERS); ONE  
 FOURTH OF THE WT CURRENT  
 (HETEROMULTIMERS).  
 FT SEQUENCE 130 AA; 14699 MW; 76717ED73C5E4D CRC64;  
 FT  
 SQ  
 Query Match 20.3%; Score 128; DB 1; Length 130;  
 Best local Similarity 41.4%; Pred. No. 5.2e-06;  
 Matches 29; Conservative 17; Mismatches 20; Indels 4; Gaps 3;  
 OY 51 LYIMWMIGSFITVALVSTVSKRREHSNDPYHOYVED--WOEKYS--QILNEESK 107  
 DB 46 LYILMVIGFEGFTLIGIMLSYIRSKLEHSHDPFNVIIESDAWQERKALFQARVLESFR 105  
 OY 108 AT-IHENIGA 116  
 DB 106 ACYVIEHQAA 115  
 DB  
 RESULT 7  
 MINK\_PIG STANDARD; PRT; 130 AA.  
 AC QSTU9;  
 DT 16-OCT-2001 (Rel. 40, Created)  
 DT 16-OCT-2001 (Rel. 40, Last sequence update)  
 DT 16-OCT-2001 (Rel. 40, Last annotation update)  
 DE Isk slow voltage-gated potassium channel protein (Minimal potassium  
 channel) (Mink).  
 GN KCNE1.  
 OS Sus scrofa (Pig).  
 OS Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Sus.  
 OX NCBI\_TaxID=9823;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Coronary artery;  
 RA Ohta S., Imaizumi Y.;  
 RA Submitted (SEP-1999) to the EMBL/Genbank/DDAJ databases.  
 CC -1- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A-POTASSIUM  
 CHANNEL. ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE  
 STABILITY OF THE MULTIMERIC COMPLEX. KCNE1 CO-ASSEMBLE WITH KCNQ1  
 (KVLQT1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC  
 POTASSIUM (IKs) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY  
 STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).  
 CC -1- SUBUNIT: ASSOCIATES WITH KCNQ1 (BY SIMILARITY).  
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.  
 CC -1- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY  
 SIMILARITY).  
 CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.  
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 CC  
 CC EMBL: AB032575; BAA86982.1; -  
 CC InterPro: IPR000369; ISK\_Channel.  
 CC Pfam: PF02060; ISK\_Channel; 1.  
 CC PRINTS: PR00168; KCNECHANNEL.  
 CC Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;

KM Glycoprotein; Phosphorylation.  
 FT DOMAIN 1 44 EXTRACELLULAR (POTENTIAL).  
 FT TRANSSEM 45 67 POTENTIAL.  
 FT DOMAIN 68 130 CYTOPLASMIC (POTENTIAL).  
 FT CARBOHYD 26 5 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT CARBOHYD 26 26 N-LINKED (GLCNAC. . .) (POTENTIAL).  
 FT MOD\_RES 103 103 PHOSPHORYLATION (BY PKC) (BY SIMILARITY).  
 FT SEQUENCE 130 AA; 14671 MW; 76717ED724E91D CRC64;  
 FT  
 SQ  
 Query Match 20.2%; Score 127.5; DB 1; Length 130;  
 Best local Similarity 42.6%; Pred. No. 5.9e-06;  
 Matches 26; Conservative 16; Mismatches 16; Indels 3; Gaps 2;  
 OY 51 LYIMWMIGSFITVALVSTVSKRREHSNDPYHOYVED--WOEKYS--QILNEESK 107  
 DB 46 LYILMVIGFEGFTLIGIMLSYIRSKLEHSHDPFNVIIESDAWQERKALFQARVLESFR 105  
 OY 108 A 108  
 DB 106 A 106  
 DB  
 RESULT 8  
 MINK\_CAVPO STANDARD; PRT; 125 AA.  
 AC Q60409; Q9QVZ5;  
 DT 30-MAY-2000 (Rel. 39, Created)  
 DT 30-MAY-2000 (Rel. 39, Last sequence update)  
 DT 16-OCT-2001 (Rel. 40, Last annotation update)  
 DE Isk slow voltage-gated potassium channel protein (Minimal potassium  
 channel) (Mink).  
 GN KCNE1.  
 OS Cavia porcellus (Guinea pig).  
 OS Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Hystriocognathi; Cavidae; Cavia.  
 OX NCBI\_TaxID=10141;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Heart;  
 RA MEDLINE=94089666; PubMed=8265583;  
 RA Varum M.D., Busch A.E., Bond C.T., Maylie J., Adelman J.P.;  
 RA "The min K channel underlies the cardiac potassium current IKs and  
 RT mediates species-specific responses to protein kinase C.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 90:11528-11532(1993).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=Heart muscle;  
 RA MEDLINE=94173910; PubMed=7510407;  
 RA Zhang J., Jurkiewicz N.K., Folander K., Lazareides E., Salata J.J.,  
 RA Swanson R.;  
 RA "K+ currents expressed from the guinea pig cardiac Isk protein are  
 RT enhanced by activators of protein kinase C.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 91:1766-1770(1994).  
 CC -1- FUNCTION: ANCILLARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM  
 CHANNEL. ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE  
 STABILITY OF THE MULTIMERIC COMPLEX. KCNE1 CO-ASSEMBLE WITH KCNQ1  
 (KVLQT1) TO FORM THE SLOWLY ACTIVATING DELAYED RECTIFIER CARDIAC  
 POTASSIUM (IKs) CHANNEL. THE OUTWARD CURRENT REACHES ITS STEADY  
 STATE ONLY AFTER 50 SECONDS (BY SIMILARITY).  
 CC -1- SUBUNIT: ASSOCIATES WITH KCNQ1 (BY SIMILARITY).  
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.  
 CC -1- PTM: PHOSPHORYLATION INHIBITS THE POTASSIUM CURRENT (BY  
 SIMILARITY).  
 CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.  
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FT CARBOND      5          5      N-LINKED (GLCNAC.. ) (POTENTIAL).
CT MOJ_RES     103        103      PHOSPHORYLATION (BY PKC) (BY SIMILARITY).
SQ SEQUENCE    130 AA; 14617 MW; E7B10048F41E205E CRC64;

Query Match           19.8%; Score 125; DB 1; Length 130;
Best Local Similarity 34.2%; Pred. No. 1e-05;
Matches 25; Conservative 20; Mismatches 16; Indels 12; Gaps 3;

OY 51 LYIWMIGMFSEITVIAIVTSFKSRRESHNDPYHOXI-VEDMDER-----YKSQILT--- 102
   ||:::| | | | | : | :: | | | | | | | | | | | | | | | | | | | | | |
DB 46 LYIIMVLGFGFFGTIGIMLSYRSKLEHSDHPFVVLYIEANDMCKDRAVYQAVYLESCR 105
   || | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |

OY 103 ----LESKATH 111
   || | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |

DB 106 GCYVENOLAVEH 118

RESULT 10
ID YZ04_METUA STANDARD: PRT; 439 AA.
YZ04_METUA
AC Q60260;
DT 01-NOV-1997 (Rel. 35, Created)
DM 01-NOV-1997 (Rel. 35, Last sequence update)
DE 16-OCT-2001 (Rel. 40, Last annotation update)
DT Hypothetical protein MJEC104.
MN MJEC104.
OS Methanococcus jannaschii.
OC Archaea; Euryarchaeota; Methanococci; Methanococcales;
CC Methanocaldococcaceae; Methanocaldococcus.
OX NCBI_TaxID=2190;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-JAL-1 / DSM 2661 / ATCC 43067;
RX MEDLINE=96337999; PubMed=8688087;
RA Bult C.J., White O., Olsen G.J., Zhou L., Fleischmann R.D.,
RA Sutton G.G., Blake J.A., Fitzgerald L.M., Clayton R.A., Goeysne J.D.,
RA Kesteven A.R., Dougherty B.A., Tomb J.-F., Adams M.D., Reich C.I.,
RA Overbeek R., Kirkness N.E.F., Weisskopf K.G., Merrick J.M., Glodek A.,
RA Scott J.L., Georgagen S.S.M., Weidman J.F., Fuhrmann J.L., Nguyen D.,
RA Uitterback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,
RA Cotton M.D., Roberts K.M., Hurst M.A., Kalne B.P., Borodovsky M.,
RA Kleck H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;
RT "Complete genome sequence of the methanogenic archaeon, Methanococcus
RT jannaschii."
RL Science 273:1058-1073(1996).
CC -I- SIMILARITY: SOME TO M.JANNASCHII MJ0425.
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CC -----
CC EMBL: L77118; AAC37078.1; -.
DR TIGR: MJEC104; -.
DR InterPro: IPR004256; DUF234.
DR InterPro: IPR004309; DUF238.
DR Pfam: PF03008; DUF234; 1.
DR Pfam: PF03075; DUF238; 1.
KW Hypothetical protein; ATP-binding; Complete proteome.
FT NP_BIND 28 35 ATP (POTENTIAL).
FT SEQUENCE 439 AA; 52421 MW; AA50918A58ERC354 CRC64;

Query Match           13.4%; Score 84.5; DB 1; Length 439;
Best Local Similarity 23.3%; Pred. No. 0.43;
Matches 24; Conservative 19; Mismatches 31; Indels 29; Gaps 3;

OY 33 EDEALQARYDAINFYYIYLMMVMGMFSFITVALIVSYVKRRRH----- 79
   |::| |::| |::| | | | | | | | | | | | | | | | | | | | | | | | | |
DB 8 ELKAINEKDSNNFEVIYIGRRIRIGKTTLAKSV-----ENNEHYIYLAEGDNLIK 60

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OY 80 -----SNDPYHOIVEDMOEKY---KSQJLNLESKATIHEN 113
DB 61 FKRYSKVEPTIEYAKEDWEAFNFKLIIIDEPNLIKEN 103

RESULT 11
CPS_CLOTH STANDARD; PRT; 557 AA.
ID CPS_CLOTH
AC 002929;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Putative sensory transducer protein.
OS Clostridium thermocellum.
OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
OC Clostridium.
OX NCBI_TaxID=1515;
RN SEQUENCE FROM N.A.
RC STRAIN=NCIB 10682;
RX MEDLINE=93171873; PubMed=8436949;
RA Hazlewood G.P., Davidson K., Laurie J.I., Huskisson N.S.,
RA Gilbert H.J.;
RT "Gene sequence and properties of cell, a family E endoglucanase from
RT Clostridium thermocellum."
RT J. Gen. Microbiol. 139:307-316(1993).
CC -1- FUNCTION: MAY BIND ATTRACTANTS OR DETECT CHANGES IN THE
CC EXTRACELLULAR CONCENTRATION OF SOLUBLE SUGARS.
CC -1- SIMILARITY: BELONGS TO THE CHEMOTAXIS SENSORY TRANSDUCERS FAMILY.
CC -----
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CC -----
DR EMBL: L04736; AAA20891.1; -.
DR PIR: B47704; B47704.
DR HSSP: P02942; 1007.
DR InterPro: IPR004089; Chmtaxis_transd.
DR InterPro: IPR003660; HAMP.
DR Pfam: PF000015; MCPsigmal; 1.
DR Pfam: PF00672; HAMP; 1.
DR SMART: SM00304; HAMP; 1.
DR SMART: SM00283; MA; 1.
DR Chemotaxis; Transducer; Transmembrane; Methylation.
DR TRANSMEM 122 145
FT MOD_RES 268 268 POTENTIAL.
FT MOD_RES 274 274 DEAMIDATION AND METHYLATION (BY
FT MOD_RES 281 281 SIMILARITY).
FT MOD_RES 281 281 METHYLATION (BY SIMILARITY).
FT MOD_RES 281 281 METHYLATION (BY SIMILARITY).
FT MOD_RES 281 281 METHYLATION (BY SIMILARITY).
SQ SEQUENCE 557 AA; 61042 MW; 169875A91F614446 CRC64;

Query Match 12.1%; Score 76.5; DB 1; Length 557;
Best Local Similarity 23.2%; Pred. No. 3.5;
Matches 29; Conservative 24; Mismatches 47; Indels 25; Gaps 5;

OY 1 MSLTSLNFTQLEDFVRIFTYMDNKRONTAEOALQAVDENFTYVILYLMWIGMF 60
DB 84 LSLDIDIGDLRYF-EAFLEY-----NTTAKKEVDENKQVASTVMIYIVFGIL 136
OY 61 SFTIIVALTSTVKRRRHSNDPYHOYI-----VE-DMQEKYSQJLNLESKAT 109
DB 137 IATLALGVFSIRISK-----PIGQVNEADRLALGDEVVDAKATRDIGKLASFPR 189
OY 110 IHENI 114
DB 190 MIENI 194

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RESULT 12
MIR2_HUMAN STANDARD; PRT; 103 AA.
ID MIR2_HUMAN
AC 091656;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Minimum potassium ion channel-related peptide 2 (MIRP2) (Mink-related
DE peptide 2).
GN KCNE3.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN SEQUENCE FROM N.A.
RA Abbott G.W., Sesti F., Buck M.R., Goldstein S.A.N.;
RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: ANCLILARY PROTEIN THAT CO-ASSEMBLE WITH A POTASSIUM
CC CHANNEL ALPHA-SUBUNIT TO MODULATE THE GATING KINETICS AND ENHANCE
CC STABILITY OF THE MULTIMERIC COMPLEX (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE KCNE FAMILY OF POTASSIUM CHANNELS.
CC -----
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CC -----
DR EMBL: AF076531; AAD28089.1; -.
DR Gene: HGNC:6243; KCNE3.
DR MIM: 604433; -.
KW Ionic channel; Transmembrane; Ion transport; Voltage-gated channel;
KW Glycoprotein.
FT DOMAIN 1 57 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 58 78 POTENTIAL.
FT DOMAIN 79 103 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 5 5 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 41 41 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 103 AA; 11710 MW; 5235385ED08BF10 CRC64;

Query Match 12.0%; Score 76; DB 1; Length 103;
Best Local Similarity 30.0%; Pred. No. 0.6;
Matches 18; Conservative 16; Mismatches 22; Indels 4; Gaps 2;

OY 29 NTAEOEALQAVDENFTYVILYLMWIGMFSFTIIVALTSTVKRRRHSNDPYHOYI 88
DB 41 NTEERRASLPGRD-DNSYVILTFVWF--LPAVVGSLILGYTRSKRKDKSDPYHYI 96

RESULT 13
SYL_BORBU STANDARD; PRT; 1042 AA.
ID SYL_BORBU
AC 051773;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Isolation-1-rRNA synthetase (EC 6.1.1.5) (Isolation-1-rRNA ligase)
DE (ILERS).
GN ILRS OR BB0833.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN SEQUENCE FROM N.A.
RA STRAIN=ATCC 35210 / B31;
RX MEDLINE=98065943; PubMed=9403685;
RA Fraser C.M., Casjens S., Huang W.M., Sutton G.G., Clayton R.A.,

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RA Lathigra R., White O., Ketchum K.A., Dodson R., Hickey E.K., Gwinn M.,
RA Dougherty B., Tomb J.-F., Fleischmann R.D., Richardson D., Hanson M.,
RA Peterson J., Kierlavage A.R., Quackenbush J., Salzberg S., Hanson M.,
RA van Vugt R., Palmer N., Adams M.D., Gocayne J.D., Weidman J.,
RA Uterback T., Matthey L., McDonald L., Artlich P., Bowman C.,
RA Garland S., Fujii C., Cotton M.D., Horst K., Roberts K., Hatch B.,
RA Smith H.O., Venter J.C.;
RT "Genomic sequence of a Lyme disease spirochaete, Borrelia
RT burgdorferi."
RT Nature 390:580-586(1997).
CC -1- CATALYTIC ACTIVITY: ATP + L-isoleucine + tRNA(ile) -> AMP +
CC diisophosphate + L-isoleucyl-tRNA(ile).
CC -1- COFACTOR: BINDS 1 ZINC ION (BY SIMILARITY).
CC -1- SUBUNIT: MONOMER (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -1- SIMILARITY: BELONGS TO CLASS-I AMINOACYL-TRNA SYNTHETASE FAMILY.
CC -----
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CC -----
DR EMBL: AE001181; AAC67179.1; -;
DR HSPB: P56690; 11LE.
DR TIGR: BB0833; -.
DR InterPro: IPR002300; tRNA-synt_1a.
DR InterPro: IPR001412; tRNA-synt_1.
DR InterPro: IPR002301; tRNA-synt_1le.
DR Pfam: PF00133; tRNA-synt_1.1.
DR PRINTS: PR00984; TRNASYNTHILE.
DR TRFAMS: TIGR00392; 1LES; 1.
DR PROSITE: PS00178; AA_TRNA_LIGASE_I; FALSE_NEG.
DR Aminoacyl-tRNA synthetase; Protein biosynthesis; Ligase; ATP-binding;
KW Metal-binding; zinc; Complete proteome.
FT SITE 48 58 "HIGH" REGION.
FT SITE 594 597 "KMSKS" REGION.
FT BINDING 597 597 ATP (BY SIMILARITY).
SQ SEQUENCE 1042 AA; 122331 MW; 6C0F7D820CA32F75 CRC64;

Query Match 11.8%; Score 74.5; DB 1; Length 1042;
Best Local Similarity 27.3%; Pred. No. 11;
Matches 33; Conservative 24; Mismatches 45; Indels 19; Gaps 8;

QY 6 NETQTLEDVFRRIFFTYMDNM-----RONTAEOBALQAKYDA-ENFYVILYLMVMIG 58
DB 703 NLTKSIESLLE--FIDKLNMMYIRSRPRPKMSND--KDKNDAYETLYAKTLMILLA 758
QY 59 MF-SFTIVAIL--VSTVAKSRKREHSD---PYHQYIVDMOEYK--SOLMLESKKATI 110
DB 759 PEPFPTTEETLYONLTKDEKDSIHLDYPRANENFTNKTEEKINLARITSMARSLR 818
QY 111 H 111
DB 819 H 819

RESULT 14
V120_HSV7J STANDARD: PRT; 938 AA.
AC PS2438;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Capsid assembly protein U30.
GN U30.
OS Human herpesvirus (type 7 / strain J1) (HHV7).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Simplexvirus.
RX NCBI_Taxid=57278;
RN [1]

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RP SEQUENCE FROM N.A.
RA Nicholas J.;
RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO FAMILY THAT GROUPS TOGETHER HSV-1 UL37,
CC EBV-1 23, EBV BOLF1, VZV 21, HSV-1 63, AND HCMV UL47.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: U43400; AAC54692.1; -.
DR Kapsid assembly.
SQ SEQUENCE 938 AA; 110170 MW; F4E39A2BF0D32BC9 CRC64;

Query Match 11.7%; Score 74; DB 1; Length 938;
Best Local Similarity 27.0%; Pred. No. 11;
Matches 30; Conservative 19; Mismatches 56; Indels 6; Gaps 4;

QY 3 TLSNFTQLEDVFRRIFFTYMDNMONTTAEQALQAKYDAENFYVILYLMVMIGRSF 62
DB 773 TLQNTIIIVERTSSNANTTYQDVNSIDCHFSNMQLQSKRNITYVIDVLTNR-NLENF 831
QY 63 ITVALIVSTKSRKREHSDPYHQY-IVEDMOEKYSOILN-ESKATIH 111
DB 832 SIASOLIEAKKLVKKO---DFYNQNVODDEFTVAKSHLNLFEKQKPTIN 879

RESULT 15
NM2M_CHOCR STANDARD: PRT; 497 AA.
AC P48903;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE MADH-ubiquitinone oxidoreductase chain 2 (Ec 1.6.5.3).
GN ND2 OR MAD2.
OS Chondrus crispus (Caragheen).
OC Mitochondrion.
OC Eukaryota; Rhodophyta; Florideophyceae; Gigartinales; Gigartinaeae;
OC Chondrus.
OX NCBI_Taxid=2769;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Apices;
RX MEDLINE=95341681; PubMed=7616569;
RA Leblanc C., Boyen C., Richard O., Bonnard G., Grienenberger J.M.,
RA Klorreg B.;
RT "Complete sequence of the mitochondrial DNA of the rhodophyte
RT Chondrus crispus (Gigartinales). Gene content and genome
RT organization."
RT J. MOL. BIOL. 250:484-495(1995).
CC -1- CATALYTIC ACTIVITY: MADH + ubiquinone -> NAD(+) + ubiquinol.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein. Mitochondrial
CC inner membrane.
CC -1- SIMILARITY: BELONGS TO THE COMPLEX I SUBUNIT 2 FAMILY.
CC -----
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CC -----
DR EMBL: Z47547; CAAB7619.1; -.
DR InterPro: IPR001750; Oxidored_q1.
DR Pfam: PF00361; oxidored_q1.1.
KW Oxidoreductase; NAD; Ubiquinone; Mitochondrion; Transmembrane.
SQ SEQUENCE 497 AA; 56853 MW; DB00BF38039F40CA CRC64;

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OM protein - protein search, using sw model

Run on: May 15, 2003, 14:22:29 ; Search time 30 Seconds  
(without alignments)  
844.793 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632

Sequence: 1 MSTLSNFTQTEDEVFRRIFI.....EESKATIHENIGAGFMKSP 123

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 671580 seqs, 206047115 residues

Total number of hits satisfying chosen parameters: 671580

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL\_21.\*  
1: sp.archaea:\*  
2: sp.bacteria:\*  
3: sp.fungi:\*  
4: sp.human:\*  
5: sp.invertebrate:\*  
6: sp.mammal:\*  
7: sp.mhc:\*  
8: sp.organelle:\*  
9: sp.phage:\*  
10: sp.plant:\*  
11: sp.potent:\*  
12: sp.virus:\*  
13: sp.vertibrate:\*  
14: sp.unclassified:\*  
15: sp.virus:\*  
16: sp.bacteriapp:\*  
17: sp.archaeap.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	560	86.6	123	11 Q9D808	Q9D808 mus musculus
2	553	87.5	123	11 Q8R127	Q8R127 mus musculus
3	354	56.0	71	6 Q9BDRO	Q9BDRO oryctolagus
4	132.5	21.0	129	11 Q912F4	Q912F4 cavia porce
5	97	15.3	43	6 Q9N100	Q9N100 sus scrofa
6	80.5	12.7	53	11 Q35438	Q35438 meriones un
7	78.5	12.4	47	11 Q9CKR9	Q9CKR9 pasteurella
8	75.5	11.9	367	16 Q9JYH3	Q9JYH3 neisseria m
9	75.5	11.9	4065	3 Q9P421	Q9P421 neurospora
10	75	11.9	1003	16 Q8YPS5	Q8YPS5 anabaena sp
11	73.5	11.6	171	12 Q69504	Q69504 human herpe
12	73.5	11.6	652	3 Q74974	Q74974 schizosacch
13	73	11.6	170	4 Q8RWG9	Q8RWG9 homo sapien
14	73	11.6	591	16 Q8YTH8	Q8YTH8 anabaena sp
15	73	11.6	1224	5 Q96209	Q96209 plasmodium
16	72.5	11.5	237	11 Q9CIV2	Q9CIV2 mus musculus

17	72.5	11.5	371	16 Q95EL2	Q95EL2 staphylococ
18	72.5	11.5	437	2 Q3587	Q3587 staphylococ
19	72.5	11.4	844	5 Q966J3	Q966J3 caenorhabd1
20	71.5	11.3	361	5 Q19083	Q19083 caenorhabd1
21	71.5	11.3	895	5 Q8T9C0	Q8T9C0 drosophila
22	71.5	11.3	899	5 Q9V7B6	Q9V7B6 drosophila
23	71.5	11.3	1013	2 Q9L9S3	Q9L9S3 streptococ
24	71.5	11.3	1386	8 Q9XMS2	Q9XMS2 tetrahymena
25	71	11.2	84	12 Q8VJ39	Q8VJ39 swinepox vi
26	71	11.2	174	16 Q8YRL6	Q8YRL6 anabaena sp
27	71	11.2	1877	10 Q9ZNM4	Q9ZNM4 sorghum bic
28	70.5	11.2	451	17 Q9VOD7	Q9VOD7 pyrococcus
29	70.5	11.2	455	17 Q8RH19	Q8RH19 pyrococcus
30	70	11.1	170	4 Q96CC4	Q96CC4 homo sapien
31	70	11.1	522	16 Q990Y7	Q990Y7 staphylococ
32	70	11.1	546	5 Q2453	Q2453 caenorhabd1
33	70	11.1	947	16 P95022	P95022 mycobacteri
34	69.5	11.1	5388	5 Q9UID0	Q9UID0 leishmania
35	69.5	11.0	188	16 Q8XJW8	Q8XJW8 clostridium
36	69.5	11.0	198	5 Q9NNV9	Q9NNV9 plasmodium
37	69.5	11.0	554	5 Q23508	Q23508 caenorhabd1
38	69.5	11.0	693	10 Q9SR87	Q9SR87 arabidopsis
39	69.5	11.0	932	3 P89499	P89499 saccharomyc
40	69.5	11.0	1154	3 Q74755	Q74755 schizosacch
41	69.5	11.0	3973	5 Q96204	Q96204 plasmodium
42	69	10.9	282	16 Q31856	Q31856 bacillus su
43	69	10.9	359	16 Q93IS3	Q93IS3 staphylococ
44	69	10.9	548	16 Q99UC9	Q99UC9 staphylococ
45	69	10.9	773	10 Q98RM5	Q98RM5 guillardia

#### ALIGNMENTS

RESULT 1

ID	Q9D808	PRELIMINARY;	PRT;	123 AA.
AC	Q9D808:			
DT	01-JUN-2001 (TREMBLrel. 17, Created)			
DT	01-JUN-2001 (TREMBLrel. 17, Last sequence update)			
DT	01-JUN-2002 (TREMBLrel. 21, Last annotation update)			
DE	2200002116R1k protein.			
GN	2200002116R1K.			
OS	Mus musculus (Mouse).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
OX	NCBI_TaxID=10090;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN=C57BL/6J; TISSUE=STOMACH;			
RX	MEDLINE=21085660; PubMed=11217851;			
RA	Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,			
RA	Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,			
RA	Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamana I.,			
RA	Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,			
RA	Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,			
RA	Fleischmann W., Gaasterland T., Gissi C., King B., Kochwa H.,			
RA	Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,			
RA	Schriml L.M., Staudt F., Suzuki R., Tomita M., Wagner U., Washio T.,			
RA	Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,			
RA	Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,			
RA	Brownstein M.J., Butt C., Fletcher C., Fujita M., Gariboldi M.,			
RA	Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,			
RA	Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombertis P.,			
RA	Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,			
RA	Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,			
RA	Suzuki H., Toyooka K., Wang K.H., Welter C., Whitaker C., Wilming L.,			
RA	Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.,			
RA	Hayashizaki Y.,			
RT	*functional annotation of a full-length mouse cDNA collection.*;			
RL	Nature 409:665-690(2001).			
DR	EMBL; AK008619; BAB25781.1; -			
DR	MGD; MGI:1916393; 2200002116R1K.			

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DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel. 1.
DR PRINTS: PR01605; KCNE2CHANNEL.
DR PRINTS: PR00168; KCNECHANNEL.
SQ SEQUENCE 123 AA; 14348 MW; 0A5DB7759BED741C CRC64;

Query Match
Best Local Similarity 84.6%; Score 560; DB 11; Length 123;
Matches 104; Conservative 14; Mismatches 5; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFRFRIFITTYMDNMRONTTAEQALQAVDAENFYVILYLMVMIGMF 60
DB 1 MATLANLTQLEDAFKKIFITTYMDSMRNTTAEQALQAVDAENFYVILYLMVMIGMF 60
QY 61 SFTIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILNDESKATIHENIGAAGFK 120
DB 61 SFTIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILNDESKATIHENIGAAGFK 120
QY 121 MSP 123
DB 121 VSP 123

RESULT 2
Q8R1Z7 PRELIMINARY; PRT; 123 AA.
AC Q8R1Z7;
DT 01-JUN-2002 (TREMBlrel. 21, Created)
DT 01-JUN-2002 (TREMBlrel. 21, Last sequence update)
DT 01-JUN-2002 (TREMBlrel. 21, Last annotation update)
DE RIKEN cDNA 2200002116 gene.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-EYE;
RA Strausberg R.;
RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL: BC022699; AAH2269.1; -.
SQ SEQUENCE 123 AA; 14370 MW; 0A52784967ED741C CRC64;

Query Match
Best Local Similarity 87.5%; Score 553; DB 11; Length 123;
Matches 103; Conservative 14; Mismatches 6; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFRFRIFITTYMDNMRONTTAEQALQAVDAENFYVILYLMVMIGMF 60
DB 1 MATLANLTQLEDAFKKIFITTYMDSMRNTTAEQALQAVDAENFYVILYLMVMIGMF 60
QY 61 SFTIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILNDESKATIHENIGAAGFK 120
DB 61 SFTIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILNDESKATIHENIGAAGFK 120
QY 121 MSP 123
DB 121 VSP 123

RESULT 3
Q9BDR0 PRELIMINARY; PRT; 71 AA.
AC Q9BDR0;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2002 (TREMBlrel. 21, Last annotation update)
DE K+/pacemaker channel beta subunit MIP1 (Fragment).
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;

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RN [1]
RP SEQUENCE FROM N.A.
RA Wyome R.T., Holmes B.A., Wyome R.S., Yu H., Wu J., Potapova I.,
RA Zuckerman J., Pan Z., Wang H., Shi W., Robinson R., El-Maghrabi R.,
RA Benjamin W., Dixon J.E., McKinnon D., Cohen I.S.;
RT Mipr1: A beta subunit for the HCN ion channel family enhances
RT expression and speeds kinetics.
RL Submitted (DEC-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF329636; AAK15527.1; -.
DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel. 1.
DR PRINTS: PR00168; KCNECHANNEL.
FT NON_TER 1 71
FT NON_TER 1 71
SQ SEQUENCE 71 AA; 8557 MW; 4A67539D4032CEA CRC64;

Query Match
Best Local Similarity 56.0%; Score 354; DB 6; Length 71;
Matches 68; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 43 AENFYVILYLMVMIGMFSEFIIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILN 102
DB 1 AENFYVILYLMVMIGMFSEFIIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILN 60
QY 103 LEESKATIHEN 113
DB 61 FEEAKATIHEN 71

RESULT 4
Q91Z94 PRELIMINARY; PRT; 129 AA.
AC Q91Z94;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-JUN-2002 (TREMBlrel. 21, Last annotation update)
DE Potassium voltage-gated channel Isk-related family member 1.
OS Cavia porcellus (Guinea pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Hystriognathi; Cavidae; Cavia.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=HEART;
RA Jiang M., Zhang M., Liu J., Tieng G.-N.;
RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AY050512; AAL13162.1; -.
DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel. 1.
DR PRINTS: PR01604; KCNE1CHANNEL.
SQ SEQUENCE 129 AA; 14645 MW; 4843171C7DC1A92A CRC64;

Query Match
Best Local Similarity 21.0%; Score 132.5; DB 11; Length 129;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

QY 51 LYLMVMIGMFSEFIIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILN 100
DB 45 LYLMVMIGMFSEFIIVALTSTVSKRREHSNDPYHOYIVEDMOEKYSQILN 95

RESULT 5
Q9N100 PRELIMINARY; PRT; 57 AA.
AC Q9N100;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-JUN-2002 (TREMBlrel. 21, Last annotation update)
DE KCNE1 (Fragment).
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.
OX NCBI_TaxID=9823;

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RN [1]
RP SEQUENCE FROM N.A.
RA Li Y., Freeman L.C.;
RT *PKC modulation of granulosa Iks does not involve a PKC site in pig
   mink.*;
RL Biophys. J. 78:208A-208A(2000).
DR EMBL: AF233358; AAF43426.1; -.
DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel.1.
DR PRINTS: PR01604; KCNE1CHANNEL.
DR PRINTS: PR00168; KCNECHANNEL.
FT NON_TER 1
FT NON_TER 57
SQ SEQUENCE 57 AA; 6726 MW; EFB66FD93473ACF6 CRC64;

Query Match
Best Local Similarity 44.6%; Score 97; DB 6; Length 57;
Matches 25; Conservative 11; Mismatches 16; Indels 4; Gaps 3;

OY 65 VALVSTVSKRREHSDPYHOYVED--WQEKYS--QILNDESKAT-IHENIGA 116
DB 2 LGIMLSYIRSKLHSHDPYNYIGDSWQEKDAVFAQVAVLENCACVYIENQAA 57

RESULT 6
OY 035438 PRELIMINARY; PRT; 43 AA.
AC 035438;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE K+ channel subunit Isk (Fragment).
GN ISK.
OS Meriones unguiculatus (Mongolian jird) (Mongolian gerbil).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Gerbillinae;
OC Meriones.
OX NCBI_TaxID=10047;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=HEART, VESTIBULAR LABYRINTH, AND STRIA VASCULARIS;
RX MEDLINE=98097798; PubMed=9435509;
RA Marcus D.C., Sunose H., Liu J., Shen Z., Scofield M.A.;
RT *P2U putative kinase receptor inhibits apical Isk/KVLQT1 channel via
   protein kinase C in vestibular dark cells.*;
RL Am. J. Physiol. 273:C2022-C2029(1997).
DR EMBL: AF029765; AAB84214.1; -.
DR InterPro: IPR000369; ISK_Channel.
DR Pfam: PF02060; ISK_Channel; 1.
KW Ionic channel.
FT NON_TER 1
FT NON_TER 43
SQ SEQUENCE 43 AA; 5062 MW; D6C5C230C87D4869 CRC64;

Query Match
Best Local Similarity 12.7%; Score 80.5; DB 11; Length 43;
Matches 19; Conservative 7; Mismatches 11; Indels 3; Gaps 2;

OY 72 VSKRREHSDPYHOYVED--WQEKYS--QILNDESKA 108
DB 2 IIRSKLHSHDPYNYIESDAQAKAVFAQVAVLENCACVYIENQAA 41

RESULT 7
OY 09CKR9 PRELIMINARY; PRT; 568 AA.
AC 09CKR9;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE HsdM.
GN HsdM OR Pm1537.
OS Pasteurella multocida.

```

```

OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
OC Pasteurella.
OX NCBI_TaxID=747;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PM70;
RX MEDLINE=21145866; PubMed=11248100;
RA May B.J., Zhang Q., Li L.L., Paustian M.L., Whitlam T.S., Kapur V.;
RT *Complete genomic sequence of Pasteurella multocida Pm70.*;
RL Proc. Natl. Acad. Sci. U.S.A. 98:3460-3465(2001).
DR EMBL: AE006190; AAK03621.1; -.
DR InterPro: IPR003665; Methylase_M.
DR InterPro: IPR002296; N12N6_mifrase.
DR InterPro: IPR003356; N6_DNA_mifrase.
DR Pfam: PF02506; Methylase_M; 1.
DR Pfam: PF02384; N6_mifrase; 1.
DR PRINTS: PR00507; N12N6MTFRASE.
KW Complete proteome.
SQ SEQUENCE 568 AA; 65503 MW; 29163A09A446C741 CRC64;

Query Match
Best Local Similarity 12.4%; Score 78.5; DB 16; Length 568;
Matches 29; Conservative 21; Mismatches 31; Indels 29; Gaps 6;

OY 23 MDNWRONTTAEQEA-----LQAKVDAENFYVYILYVMIGNFSEFVAILVSTVKS 74
DB 1 MNSEQOYLNEIDAKLMKSADRLRSNIEAANKVHVLISLFL---KVSADFLA----- 51

OY 75 KRREHS-----NDPYHOYVE-----DMQEKYSQILN-LEESKATIHENI 114
DB 52 --RHSIQOQLDPEHLTYLDPSEFYDSEEQOALANELDLDYTYEENV 99

RESULT 8
OY 09UYH3 PRELIMINARY; PRT; 367 AA.
AC 09UYH3;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE Protease, putative.
GN NMB1587.
OS Neisseria meningitidis (serogroup B).
OC Bacteria; Proteobacteria; beta subdivision; Neisseriaceae; Neisseria.
OX NCBI_TaxID=491;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MC58 / SEROGROUP B;
RX MEDLINE=20175755; PubMed=10710307;
RA Tettelin H., Saunders N.J., Heidelberg J., Jeffries A.C., Nelson K.E.,
RA Eisen J.A., Ketchum K.A., Hood D.W., Peden J.F., Dodson R.J.,
RA Nelson W.C., Gwinn M.L., DeBoy R., Peterson J.D., Hickey E.K.,
RA Haft D.H., Salzberg S.L., White O., Fleischmann R.D., Dougherty B.A.,
RA Mason T., Ciecko A., Parksey D.S., Blair E., Citron H., Clark E.B.,
RA Cotton M.D., Ullrichback T.R., Khouli H., Qin H., Yamaharan J.,
RA Gill J., Scarlato V., Masiagnani V., Pizsa M., Grandi G., Sun L.,
RA Smith H.O., Fraser C.M., Moxon E.R., Rappunoli R., Venter J.C.;
RT *Complete genome sequence of Neisseria meningitidis serogroup B strain
   MC58.*;
RL Science 287:1809-1815(2000).
DR EMBL: AE002509; AAF41940.1; -.
DR MEROPS: S49.002; -.
DR TIGR: NMB1587; -.
DR InterPro: IPR002142; Peptidase_U7.
DR Pfam: PF01343; Peptidase_U7; 1.
DR ProDom: PD002897; Peptidase_U7; 1.
KW Complete proteome.
SQ SEQUENCE 367 AA; 41414 MW; 814711B592C6684 CRC64;

Query Match
Best Local Similarity 11.9%; Score 75.5; DB 16; Length 367;
Matches 25; Conservative 17; Mismatches 25; Indels 17; Gaps 4;

```

```
QY      33 EELAA--VVDADENYYVYLKAMIMGISEPIIVALLSVYSKRREHNDPHYIYE 90
Dd      11 EQEISGMKEILLNGIFLELTVFQAALIVAI---VSKKOSSESG---SVILF 62
QY      91 DMOEKIKSQ-----ITNLESK 107
Dd      63 DFSENYYKKROSFEAFFLSGEFAK 86
```

## RESULT 9

ID Q9P4Z1 PRELIMINARY; PRT; 4065 AA.

DT 01-OCT-2000 (TREMBLrel. 15, Created)  
 DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)  
 DE Related to TOM1 protein.  
 GN B11B22.010.  
 OS Neurospora crassa.  
 OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;  
 OC Sordariales; Sordariaceae; Neurospora.  
 NCBI\_TaxID=5141;

SQ SEQUENCE 4065 AA; 452568 MW; F74683CEC36F9350 CRC64;

QY 6 NFOUJLEUVFRIETTYDNNRQNTTAQDEALQAVDANFNFYVILLYLMWJMGSSFTV 65  
Db 2937 NAEQOAEERPRRYVITLKNQ-----QIDITLQIDE-----YELALPEEFDEVI 2982  
QY 66 AILVSTVSKRRH-----SNDPYHQYIVEDWQEKYQQLINLESKATIHENIGANG 118  
2963 AQAISTRSQAREOVSCGEENTFVQDELEALPEELRNILILQEOHQEORRRRONANG 3040

## RESULT 10

ID Q8YPT5 PRELIMINARY; PRT; 1003 AA.

DT 01-MAR-2002 (TREMblrel. 20, Created)  
DT 01-MAR-2002 (TREMblrel. 20, Last sequence update)  
DT 01-JUN-2002 (TREMblrel. 21, Last annotation update)  
DE Toxin secretion ABC transporter ATP-binding protein  
DE ALR4239.

BL DNA Res: 8:205-213(2001).  
DR EMEL, AP003595; BAE75938.1. --  
DR InterPro: IPR003593; AAA\_AtPase.  
DR InterPro: IPR001440; ABCtransp1TW.  
DR InterPro: IPR003439; ABC\_transport.  
DR InterPro: IPR000595; CNMP\_binding.  
DR InterPro: IPR001950; TIF-SUI1.  
DR Pfam: PF00664; ABC\_membrane.1.  
DR Pfam: PF00005; ABC\_tran.1.  
DR Pfam: PF00027; CNMP\_binding.1.  
DR Prodom: PD000006; ABC\_transportr.1.  
DR SMART: SM00382; AAA.1.  
DR SMART: SM00100; CNMP.1.  
DR PROSITE: PS00211; ABC\_TRANSPORTER.1.  
DR PROSITE: PS00042; CNMP\_BINDING.3.1.  
DR PROSITE: PS01118; SUI1.1; UNKNOWN.1.  
FW ATP-binding; Complete proteome.  
SQ SEQUENCE 1003 AA; 111718 MW; 1260643429DC6CFA CFC64;

Query Match	11.98;	Score 75;	DB 16;	Length 1003;
Best Local Similarity	23.08;	Pred. No. 30;		
Matches 31; Conservative	20;	Mismatches 30;	Indels 54;	Gaps 7;

```

OY      20 ITYDWMQONTAQBDAOLQAVDKNFYVYLYLTMV-----IGMS---FIIVAI 67
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      548 INELENIQFLEGT--TALTAVLDA--VFSSVYIIVALEFSMQLTVGLGTFPIFIITL 602
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :
OY      68 LVSYVSKSR---REHSNDPIHQIYVE-----DMQETKSOILNLEE 105
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      603 IASPSVSKQLSKSRKRNSETSYLVEWMSGIQTVKAONIELSRFSMOERIARY----- 657
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :
OY      106 SKATIHENIGAAGFK 120
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      658 -----AAGFK 662
           : : : : : : : : : : : : : : : : : : : : : : : : : : : :

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## RESULT 11

ID	Q69504	PRELIMINARY;	PRT;	171	AA.
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AC Q59504;  
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)  
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)  
 DT 01-MAR-2002 (TrEMBLrel. 20, Last annotation update)  
 DE U23 protein.  
 GN U23.  
 OS Human herpesvirus (Type 7 / strain J1) (HHV7).  
 OS Viruses; dsDNA viruses, no RNA stage; Herpesviridae  
 OC Alphaherpesvirinae; Simplexvirus.  
 NCBI\_TaxID=57278;

Query Match	11.6%	Score 73.5	DB 12	Length 171
Best Local Similarity	36.6%	Pred. 6.3		
Matches 15; Conservative	11	Mismatches	12	Gaps 1

50 ILYMMIGMFSFI--IVAILVSTVKSRRHSNDPYHÖY 87



Db 121 VLFLVILIAVAFSIGIVALALVLLIKNPKRNHKKPKRMAY 161

## RESULT 12

074974 PRELIMINARY; PRT; 652 AA.  
 AC 074974;  
 DT 01-NOV-1998 (TREMBLrel. 08, Created)  
 DT 01-NOV-1998 (TREMBLrel. 08, last sequence update)  
 DT 01-DEC-2001 (TREMBLrel. 19, last annotation update)  
 DE Hypothetical 75.7 kDa protein.  
 GN SPCC1827.01C.  
 OS Schizosaccharomyces pombe (fission yeast).  
 OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;  
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;  
 OC Schizosaccharomyces.  
 NC NCB1\_TaxID=4896;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-972H-;  
 RA Seeger K., Harris D., Wood V., Rajandream M.A., Barrell B.G.;  
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AL023777; CAI9309.1; -  
 KW Hypothetical protein.  
 SQ SEQUENCE 652 AA; 75651 MW; 96BB732F2273DE6B CRC64;

Query Match 11.6%; Score 73.5; DB 3; Length 652;  
 Best Local Similarity 22.0%; Pred. No. 27;  
 Matches 22; Conservative 19; Mismatches 30; Indels 29; Gaps 2;

QY 23 MNMWRONTAEOALQAKYDAENFYVILYLMWIGMFSFIIVALLSVYKSKRREHNSD 82  
 ID 08WVG9 PRELIMINARY; PRT; 170 AA.  
 AC 08WVG9;  
 DT 01-MAR-2002 (TREMBLrel. 20, Created)  
 DT 01-MAR-2002 (TREMBLrel. 20, last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, last annotation update)  
 DE Potassium voltage-gated channel-like protein.  
 GN KCNE4.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 NC NCB1\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-KIDNEY, AND HEART;  
 RA Hul R., Teng S., Lin C., Ma L., Zhen Y.;  
 RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AY065987; AAL49979.1; -  
 DR InterPro: IPR00369; ISK\_Channel.  
 DR PRINTS: PRO0168; KCNECHANNEL.  
 SQ SEQUENCE 170 AA; 18397 MW; 1C6FBCF87298F6C0 CRC64;

## RESULT 13

08WVG9 PRELIMINARY; PRT; 170 AA.  
 AC 08WVG9;  
 DT 01-MAR-2002 (TREMBLrel. 20, Created)  
 DT 01-MAR-2002 (TREMBLrel. 20, last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, last annotation update)  
 DE Potassium voltage-gated channel-like protein.  
 GN KCNE4.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 NC NCB1\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-KIDNEY, AND HEART;  
 RA Hul R., Teng S., Lin C., Ma L., Zhen Y.;  
 RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AY065987; AAL49979.1; -  
 DR InterPro: IPR00369; ISK\_Channel.  
 DR PRINTS: PRO0168; KCNECHANNEL.  
 SQ SEQUENCE 170 AA; 18397 MW; 1C6FBCF87298F6C0 CRC64;

Query Match 11.6%; Score 73; DB 4; Length 170;  
 Best Local Similarity 27.7%; Pred. No. 7.1;  
 Matches 23; Conservative 16; Mismatches 34; Indels 10; Gaps 3;

QY 23 MNMWRONTAEOALQAKYDA-----ENFYVILYLMWIGMFSFIIVALLSVYKSKR 77  
 ID 08WVG9 PRELIMINARY; PRT; 170 AA.  
 AC 08WVG9;  
 DT 01-MAR-2002 (TREMBLrel. 20, Created)  
 DT 01-MAR-2002 (TREMBLrel. 20, last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, last annotation update)  
 DE Potassium voltage-gated channel-like protein.  
 GN KCNE4.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 NC NCB1\_TaxID=9606;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-KIDNEY, AND HEART;  
 RA Hul R., Teng S., Lin C., Ma L., Zhen Y.;  
 RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AY065987; AAL49979.1; -  
 DR InterPro: IPR00369; ISK\_Channel.  
 DR PRINTS: PRO0168; KCNECHANNEL.  
 SQ SEQUENCE 170 AA; 18397 MW; 1C6FBCF87298F6C0 CRC64;

Db 64 EKKSSLLILYRDERLWGEAMKS 86

## RESULT 14

08YTH8 PRELIMINARY; PRT; 591 AA.  
 AC 08YTH8;  
 DT 01-MAR-2002 (TREMBLrel. 20, Created)  
 DT 01-MAR-2002 (TREMBLrel. 20, last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, last annotation update)  
 DE Two-component sensor histidine kinase.  
 GN ALR2739.  
 OS Anabaena sp. (strain PCC 7120).  
 OC Bacteria; Cyanobacteria; Nostocales; Nostocaceae; Nostoc.  
 NC NCB1\_TaxID=103690;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=21595285; PubMed=11759840;  
 RA Kaneko T., Nakamura Y., Wolk C.P., Kuritz T., Sasamoto S.,  
 RA Watanabe A., Iriguchi M., Ishikawa A., Kawashima K., Kimura T.,  
 RA Kishida Y., Kohara M., Matsumoto M., Matsuno A., Muraki A.,  
 RA Nakazaki N., Shimp S., Sugimoto M., Takazawa M., Yamada M.,  
 RA Yasuda M., Tabata S.;  
 RT Complete genomic sequence of the filamentous nitrogen-fixing  
 cyanobacterium Anabaena sp. strain PCC 7120.\*;  
 RL DNA Res. 8:205-213(2001).  
 DR EMBL; AP003590; BAB74438.1; -  
 DR InterPro: IPR003594; ATPbind\_ATPase.  
 DR InterPro: IPR004358; Bact\_sens\_pr\_C.  
 DR InterPro: IPR003660; HAMP.  
 DR InterPro: IPR004359; HIS\_KIN\_sig.  
 DR Pfam: PF00672; HAMP; 1.  
 DR Pfam: PF02518; HATPase\_C; 1.  
 DR Pfam: PF00512; signal; 1.  
 DR PRINTS: PR00344; BCTRLSENSOR.  
 DR SMART: SMO0304; HAMP; 1.  
 DR SMART: SMO0387; HATPase\_C; 1.  
 DR SMART: SMO0388; HASKA; 1.  
 DR PROSITE: PS0109; HIS\_KIN; 1.  
 KW Kinase; Complete proteome.  
 SQ SEQUENCE 591 AA; 67107 MW; B328BB6D2696C5D2 CRC64;

Query Match 11.6%; Score 73; DB 16; Length 591;  
 Best Local Similarity 21.3%; Pred. No. 28;  
 Matches 30; Conservative 22; Mismatches 35; Indels 54; Gaps 6;

QY 4 LSNFTQT---LEDYFRIRIFTYMDNWRONTAEOALQAKYDAENFY-----YVI 50  
 ID 08YTH8 PRELIMINARY; PRT; 591 AA.  
 AC 08YTH8;  
 DT 01-MAR-2002 (TREMBLrel. 20, Created)  
 DT 01-MAR-2002 (TREMBLrel. 20, last sequence update)  
 DT 01-JUN-2002 (TREMBLrel. 21, last annotation update)  
 DE Two-component sensor histidine kinase.  
 GN ALR2739.  
 OS Anabaena sp. (strain PCC 7120).  
 OC Bacteria; Cyanobacteria; Nostocales; Nostocaceae; Nostoc.  
 NC NCB1\_TaxID=103690;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE=21595285; PubMed=11759840;  
 RA Kaneko T., Nakamura Y., Wolk C.P., Kuritz T., Sasamoto S.,  
 RA Watanabe A., Iriguchi M., Ishikawa A., Kawashima K., Kimura T.,  
 RA Kishida Y., Kohara M., Matsumoto M., Matsuno A., Muraki A.,  
 RA Nakazaki N., Shimp S., Sugimoto M., Takazawa M., Yamada M.,  
 RA Yasuda M., Tabata S.;  
 RT Complete genomic sequence of the filamentous nitrogen-fixing  
 cyanobacterium Anabaena sp. strain PCC 7120.\*;  
 RL DNA Res. 8:205-213(2001).  
 DR EMBL; AP003590; BAB74438.1; -  
 DR InterPro: IPR003594; ATPbind\_ATPase.  
 DR InterPro: IPR004358; Bact\_sens\_pr\_C.  
 DR InterPro: IPR003660; HAMP.  
 DR InterPro: IPR004359; HIS\_KIN\_sig.  
 DR Pfam: PF00672; HAMP; 1.  
 DR Pfam: PF02518; HATPase\_C; 1.  
 DR Pfam: PF00512; signal; 1.  
 DR PRINTS: PR00344; BCTRLSENSOR.  
 DR SMART: SMO0304; HAMP; 1.  
 DR SMART: SMO0387; HATPase\_C; 1.  
 DR SMART: SMO0388; HASKA; 1.  
 DR PROSITE: PS0109; HIS\_KIN; 1.  
 KW Kinase; Complete proteome.  
 SQ SEQUENCE 591 AA; 67107 MW; B328BB6D2696C5D2 CRC64;

## RESULT 15

096209 PRELIMINARY; PRT; 1224 AA.  
 AC 096209;  
 DT 01-MAY-1999 (TREMBLrel. 10, Created)  
 DT 01-MAY-1999 (TREMBLrel. 10, last sequence update)  
 DT 01-MAY-1999 (TREMBLrel. 10, last annotation update)  
 DE Hypothetical 144.0 kDa protein.  
 GN PF05080W.  
 OS Plasmodium falciparum.  
 OC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.  
 NC NCB1\_TaxID=5833;  
 RN [1]

RP SEQUENCE FROM N.A.  
RX MEDLINE-99021743; PubMed-9804551;  
RA Gardner M.J., Tetteila H., Carucci D.J., Cummings L.M., Aravind L.,  
RA Koonin E.V., Shallow S., Mason T., Yu K., Fujii C., Pederson J.,  
RA Shen K., Jing J., Aston C., Lal Z., Schwartz D.C., Perlea M.,  
RA Salzberg S., Zhou L., Sutton G.G., Clayton R., White O., Smith R.O.,  
RA Fraser C.M., Adams M.D., Venter J.C., Hoffman S.L.,  
RT "Chromosome 2 sequence of the human malaria parasite Plasmodium  
falciparum";  
RL Science 282:1126-1132(1998).  
DR EMBL: AE001404; AAC71905.1; -.  
KW Hypothetical protein.  
SQ SEQUENCE 1224 AA; 144031 MW; B585BB28D47BDC3F CRC64;

Query Match 11.6%; Score 73; DB 5; Length 1224;  
Best Local Similarity 18.6%; Pred. No. 61;  
Matches 13; Conservative 23; Mismatches 34; Indels 0; Gaps 0;

QY 36 ALQAKVDAENFYVYLVMIGWSEFTIVAILVSTVSKRRRHSNDPYHQYIVEDWQEK 95  
DB 1001 SVAAVYQINNPLNLYVYLSNSYGIILAKLIGYISSQKRREKDNQNKRYEYNILKEPMKE 1060  
QY 96 YKSGILNLEE 105  
DB 1061 YTKLFVEKNE 1070

Search completed: May 15, 2003, 14:26:13  
Job time : 34 secs

GenCore version 5.1.4.p5\_4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 15, 2003, 14:25:44 ; Search time 53 Seconds  
(without alignments)  
223.839 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632

Sequence: 1 MSTLSNFTQLEDFVRRIRFI.....ESKATIHENIGAGFKMSP 123

Scoring table: BLOSUM62

Searched: 362588 seqs, 96450795 residues

Total number of hits satisfying chosen parameters: 362588

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Published\_Applications\_AA:\*  
1: /cgn2\_6/ptodata/2/pubpaa/US08\_NEM\_PUB pep:\*  
2: /cgn2\_6/ptodata/2/pubpaa/PCr\_NEM\_PUB pep:\*  
3: /cgn2\_6/ptodata/2/pubpaa/US06\_NEM\_PUB pep:\*  
4: /cgn2\_6/ptodata/2/pubpaa/US06\_PUBCOMB pep:\*  
5: /cgn2\_6/ptodata/2/pubpaa/US07\_NEM\_PUB pep:\*  
6: /cgn2\_6/ptodata/2/pubpaa/US07\_PUBCOMB pep:\*  
7: /cgn2\_6/ptodata/2/pubpaa/PCrUS\_PUBCOMB pep:\*  
8: /cgn2\_6/ptodata/2/pubpaa/US08\_PUBCOMB pep:\*  
9: /cgn2\_6/ptodata/2/pubpaa/US09\_NEM\_PUB pep:\*  
10: /cgn2\_6/ptodata/2/pubpaa/US09\_PUBCOMB pep:\*  
11: /cgn2\_6/ptodata/2/pubpaa/US10\_NEM\_PUB pep:\*  
12: /cgn2\_6/ptodata/2/pubpaa/US10\_PUBCOMB pep:\*  
13: /cgn2\_6/ptodata/2/pubpaa/US60\_NEM\_PUB pep:\*  
14: /cgn2\_6/ptodata/2/pubpaa/US60\_PUBCOMB pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	632	100.0	123	9	US-10-000-151B-4
2	632	100.0	123	9	US-09-864-761-49007
3	533	84.3	103	10	US-09-864-761-36713
4	132.5	21.0	76	10	US-09-864-761-37234
5	132.5	21.0	129	9	US-10-138-316-4
6	70	11.1	523	10	US-09-815-242-5723
7	70	11.1	525	10	US-09-815-242-12651
8	68.5	10.8	1590	9	US-10-180-326-1
9	67.5	10.7	380	10	US-09-134-333-12
10	67	10.6	272	10	US-09-815-242-5572
11	67	10.6	277	10	US-09-815-242-1282
12	66.5	10.5	380	10	US-09-134-333-13
13	65.5	10.4	553	10	US-09-815-242-5778
14	65.5	10.4	1276	10	US-09-982-610-24
15	65.5	10.4	2016	9	US-09-866-994-2
16	65.5	10.4	2016	10	US-09-840-125-4
17	65	10.3	367	10	US-09-815-242-10676
18	65	10.3	1482	10	US-09-815-242-12484
19	65	10.3	1835	10	US-09-935-541-5

20	65	10.3	2175	10	US-09-935-541-2	Sequence 2, Appl1
21	65	10.3	2188	10	US-09-935-541-4	Sequence 4, Appl1
22	64.5	10.2	425	9	US-10-174-590-570	Sequence 570, App
23	64.5	10.2	425	9	US-10-176-758-570	Sequence 570, App
24	64.5	10.2	425	9	US-10-175-737-570	Sequence 570, App
25	64.5	10.2	425	9	US-10-173-706-570	Sequence 570, App
26	64.5	10.2	425	9	US-10-175-738-570	Sequence 570, App
27	64.5	10.2	425	9	US-10-175-752-570	Sequence 570, App
28	64.5	10.2	425	9	US-10-176-482-570	Sequence 570, App
29	64.5	10.2	425	9	US-10-176-757-570	Sequence 570, App
30	64.5	10.2	425	9	US-10-176-913-570	Sequence 570, App
31	64.5	10.2	425	9	US-10-180-552-570	Sequence 570, App
32	64.5	10.2	425	9	US-10-180-557-570	Sequence 570, App
33	64.5	10.2	425	9	US-10-173-700-570	Sequence 570, App
34	64.5	10.2	425	9	US-10-174-572-570	Sequence 570, App
35	64.5	10.2	425	9	US-10-174-579-570	Sequence 570, App
36	64.5	10.2	425	9	US-10-174-582-570	Sequence 570, App
37	64.5	10.2	425	9	US-10-174-588-570	Sequence 570, App
38	64.5	10.2	425	9	US-10-175-739-570	Sequence 570, App
39	64.5	10.2	425	9	US-10-175-740-570	Sequence 570, App
40	64.5	10.2	425	9	US-10-175-743-570	Sequence 570, App
41	64.5	10.2	425	9	US-10-176-488-570	Sequence 570, App
42	64.5	10.2	425	9	US-10-176-492-570	Sequence 570, App
43	64.5	10.2	425	9	US-10-176-747-570	Sequence 570, App
44	64.5	10.2	425	9	US-10-176-750-570	Sequence 570, App
45	64.5	10.2	425	9	US-10-176-985-570	Sequence 570, App

ALIGNMENTS

RESULT 1  
US-10-000-151B-4  
; Sequence 4, Application US/10000151B  
; Publication No. US20030013136A1  
; GENERAL INFORMATION:  
; APPLICANT: Balseer, Jeffrey R.  
; TITLE OF INVENTION: HUMAN KCR1 REGULATION OF HERG POTASSIUM CHANNEL BLOCK  
; FILE REFERENCE: Vanderbilt Ref No. US20030013136A1 V00120; Attorney Docket No. US2  
; CURRENT FILING DATE: 2000-10-30  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 4  
; LENGTH: 123  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; US-10-000-151B-4

Query Match 100.0%; Score 632; DB 9; Length 123;  
Best Local Similarity 100.0%; Pred. No. 3.9e-61;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFVRRIRFIITMDNMKRONTTAEQALQAKYDAENFYVILYIMGMF 60  
DB 1 MSTLSNFTQLEDFVRRIRFIITMDNMKRONTTAEQALQAKYDAENFYVILYIMGMF 60  
QY 61 SFTIIVLVSTYVSKRRHSNDPYHQYIVEDWQEKYSQILNLESKATIHENIGAGFK 120  
DB 61 SFTIIVLVSTYVSKRRHSNDPYHQYIVEDWQEKYSQILNLESKATIHENIGAGFK 120  
QY 121 MSP 123  
DB 121 MSP 123

RESULT 2  
US-09-864-761-49007  
; Sequence 49007, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharron G.

```

1 APPLICANT: Rank, David R.
2 APPLICANT: Hanzel, David K.
3 APPLICANT: Chen, Wensheng
4 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
5 TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
6 FILE REFERENCE: Aeo mica-x-1
7 CURRENT APPLICATION NUMBER: US/09/864,761
8 CURRENT FILING DATE: 2001-05-23
9 PRIOR APPLICATION NUMBER: US 60/180,312
10 PRIOR FILING DATE: 2000-02-04
11 PRIOR APPLICATION NUMBER: US 60/207,456
12 PRIOR FILING DATE: 2000-05-26
13 PRIOR APPLICATION NUMBER: US 09/632,366
14 PRIOR FILING DATE: 2000-08-03
15 PRIOR APPLICATION NUMBER: GB 24263.6
16 PRIOR FILING DATE: 2000-10-04
17 PRIOR APPLICATION NUMBER: US 60/236,359
18 PRIOR FILING DATE: 2000-09-27
19 PRIOR APPLICATION NUMBER: PCT/US01/00666
20 PRIOR FILING DATE: 2001-01-30
21 PRIOR APPLICATION NUMBER: PCT/US01/00667
22 PRIOR FILING DATE: 2001-01-30
23 PRIOR APPLICATION NUMBER: PCT/US01/00664
24 PRIOR FILING DATE: 2001-01-30
25 PRIOR APPLICATION NUMBER: PCT/US01/00669
26 PRIOR FILING DATE: 2001-01-30
27 PRIOR APPLICATION NUMBER: PCT/US01/00665
28 PRIOR FILING DATE: 2001-01-30
29 PRIOR APPLICATION NUMBER: PCT/US01/00668
30 PRIOR FILING DATE: 2001-01-30
31 PRIOR APPLICATION NUMBER: PCT/US01/00663
32 PRIOR FILING DATE: 2001-01-30
33 PRIOR APPLICATION NUMBER: PCT/US01/00662
34 PRIOR FILING DATE: 2001-01-30
35 PRIOR APPLICATION NUMBER: PCT/US01/00661
36 PRIOR FILING DATE: 2001-01-30
37 PRIOR APPLICATION NUMBER: PCT/US01/00670
38 PRIOR FILING DATE: 2001-01-30
39 PRIOR APPLICATION NUMBER: US 60/234,687
40 PRIOR FILING DATE: 2000-09-21
41 PRIOR APPLICATION NUMBER: US 09/608,408
42 PRIOR FILING DATE: 2000-06-30
43 PRIOR APPLICATION NUMBER: US 09/774,203
44 PRIOR FILING DATE: 2001-01-29
45 NUMBER OF SEQ ID NOS: 49117
46 SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
47 SEQ ID NO 49007
48 LENGTH: 123
49 TYPE: PRT
50 ORGANISM: Homo sapiens
51 FEATURE:
52 OTHER INFORMATION: MAP TO AP000120.1
53 OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.98
54 OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 0.67
55 OTHER INFORMATION: EST_HUMAN HIT: A1962650.1, EVALU6 3.00e-59
56 OTHER INFORMATION: SWISSPROT HIT: Q9Y656, EVALU6 7.00e-67
57 US-09-864-761-49007

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RESULT 3
US-09-864-761-36713
: Sequence 36713, Application US/09864761
: Patent No. US20020048763A1
GENERAL INFORMATION:
: APPLICANT: Penn, Sharron G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263,6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
SEQ ID NO 36713
LENGTH: 103
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000052.1
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
OTHER INFORMATION: EXPRESSED IN HBLL00, SIGNAL = 1.1
OTHER INFORMATION: EXPRESSED IN ADUL LIVER, SIGNAL = 1.3
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.92
OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.4
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.2
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
OTHER INFORMATION: SWISSPROT HIT: Q916U6, EVALU6 2.00e-55
OTHER INFORMATION: EST_HUMAN HIT: A1962650.1, EVALU6 1.00e-54
US-09-864-761-36713

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Query Match 84.3%; Score 533; DB 10; Length 103;  
Best Local Similarity 100.0%; Pred. No. 1.7e-50;  
Matches 103; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 19 FTYYDMNRONTAEOALQAFVDAENFYVILYLMVMIGMSFTIIVASTVSKRRE 78  
DB 1 FTYYDMNRONTAEOALQAFVDAENFYVILYLMVMIGMSFTIIVASTVSKRRE 60  
QY 79 HNDPYHOYIVEDMOEKYSQILNEESKATIHENIGAGFKM 121  
DB 61 HNDPYHOYIVEDMOEKYSQILNEESKATIHENIGAGFKM 103

RESULT 4  
US-09-864-761-37234

; Sequence 37234, Application US/09864761  
; Patent No. US2002048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharon G.  
; APPLICANT: Rank, David R.  
; APPLICANT: Hanzel, David K.  
; APPLICANT: Chen, Wensheng  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: Aecm1ca-X-1  
; CURRENT APPLICATION NUMBER: US/09/864,761

; CURRENT FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/180,312  
; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 09/632,366  
; PRIOR FILING DATE: 2000-08-03  
; PRIOR APPLICATION NUMBER: GB 24263.6  
; PRIOR FILING DATE: 2000-10-04  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; PRIOR APPLICATION NUMBER: PCT/US01/00666  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00662  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00661  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/00670  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 60/234,687  
; PRIOR FILING DATE: 2000-09-21  
; PRIOR APPLICATION NUMBER: US 09/608,408  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: US 09/774,203  
; PRIOR FILING DATE: 2001-01-29  
; NUMBER OF SEQ ID NOS: 49117  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 37234

; LENGTH: 76  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AF000121.1  
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL - 1.1

; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL - 0.94  
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL - 0.59  
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL - 0.74  
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL - 0.66  
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL - 0.66  
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL - 0.7  
; OTHER INFORMATION: EST HUMAN HIT: A1246239.1, EVALUATE 2.00e-07  
; OTHER INFORMATION: SWISSPROT HIT: P15382, EVALUATE 1.00e-39  
US-09-864-761-37234

Query Match 21.0%; Score 132.5; DB 10; Length 76;  
Best Local Similarity 45.1%; Pred. No. 2.8e-07;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

QY 51 LYLMVMIGMSFTIIVASTVSKRREHSDPYHOYIVED-MOEKYSQI 100  
DB 9 LYLMVIGFEGFETGIMLSYRSKLEHSDPNVYIESDAMOEKDKAVY 59

RESULT 5  
US-10-138-316-4

; Sequence 4, Application US/10138316  
; Publication No. US20030054380A1  
; GENERAL INFORMATION:  
; APPLICANT: Keating, Mark T.  
; APPLICANT: Sanguinetti, Michael C.  
; APPLICANT: Spiawski, Igor  
; TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MINK WHICH  
; TITLE OF INVENTION: CAUSE ARRYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING  
; FILE REFERENCE: 2323-162  
; CURRENT APPLICATION NUMBER: US/10/138,316

; CURRENT FILING DATE: 2002-05-06  
; PRIOR APPLICATION NUMBER: 09/444,295  
; PRIOR FILING DATE: 1999-11-22  
; PRIOR APPLICATION NUMBER: 09/135,020  
; PRIOR FILING DATE: 1998-08-17  
; PRIOR APPLICATION NUMBER: 08/921,068  
; PRIOR FILING DATE: 1997-08-29  
; PRIOR APPLICATION NUMBER: 08/739,383  
; PRIOR FILING DATE: 1996-10-29  
; PRIOR APPLICATION NUMBER: 60/019,014  
; PRIOR FILING DATE: 1995-12-22  
; PRIOR APPLICATION NUMBER: 60/094,477  
; PRIOR FILING DATE: 1998-07-29  
; NUMBER OF SEQ ID NOS: 114  
; SOFTWARE: PatentIn ver. 2.0  
; SEQ ID NO 4  
; LENGTH: 129  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
US-10-138-316-4

Query Match 21.0%; Score 132.5; DB 9; Length 129;  
Best Local Similarity 45.1%; Pred. No. 5.4e-07;  
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

QY 51 LYLMVMIGMSFTIIVASTVSKRREHSDPYHOYIVED-MOEKYSQI 100  
DB 45 LYLMVIGFEGFETGIMLSYRSKLEHSDPNVYIESDAMOEKDKAVY 95

RESULT 6  
US-09-815-242-5723

; Sequence 5723, Application US/09815242  
; Patent No. US20020061569A1  
; GENERAL INFORMATION:  
; APPLICANT: Haselbeck, Robert  
; APPLICANT: Ohlsen, Karl L.  
; APPLICANT: Zyskind, Judith W.  
; APPLICANT: Wall, Daniel  
; APPLICANT: Trawick, John D.  
; APPLICANT: Carr, Grant J.

```

? APPLICANT: Yamamoto, Robert T.
? APPLICANT: Xu, H. Howard
? TITLE OF INVENTION: Identification of Essential Genes in
? TITLE OF INVENTION: Prokaryotes
? FILE REFERENCE: ELITRA.011A
? CURRENT APPLICATION NUMBER: US/09/915.242
? PRIOR FILING DATE: 2001-03-21
? PRIOR APPLICATION NUMBER: 60/191,078
? PRIOR FILING DATE: 2000-03-21
? PRIOR APPLICATION NUMBER: 60/206,848
? PRIOR FILING DATE: 2000-05-23
? PRIOR APPLICATION NUMBER: 60/207,727
? PRIOR FILING DATE: 2000-05-26
? PRIOR APPLICATION NUMBER: 60/242,578
? PRIOR FILING DATE: 2000-10-23
? PRIOR APPLICATION NUMBER: 60/253,625
? PRIOR FILING DATE: 2000-11-27
? PRIOR APPLICATION NUMBER: 60/257,931
? PRIOR FILING DATE: 2000-12-22
? PRIOR APPLICATION NUMBER: 60/269,308
? PRIOR FILING DATE: 2001-02-16
? NUMBER OF SEQ ID NOS: 1410
? SOFTWARE: FastSeq for Windows Version 4.0
? SEQ ID NO 5723
? LENGTH: 523
? TYPE: PRF
? ORGANISM: Staphylococcus aureus
? US-09-815-242-5723

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Query Match      11.1%; Score 70; DB 10; Length 523;
Best Local Similarity 27.9%; Pred. No. 18;
Matches 29; Conservative 14; Mismatches 33; Indels 28; Gaps 5;

QY      35  EALQAKVDANFYYVILYLMWIGFSEFLIYALVSTVSKRKRHSNDPHYVED-WQ 93
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Db      44  EATRIAYEAE-----GKFDIYYIOAPYSEFLTNILQMISEPIYTVDESEWS 90

QY      94  -----EKYKSQL---NLEE--SKATIHENIGAGFKKSP 123
      :|| | | | | | | | | | | | | | | | | | | | |
Db      91  VEYEDENVQKTYVQPLHYRNIIEENNNKLEAVSESGYGQDKVSP 134

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1      RESULT 7
2      US-09-815-242-12651
3      ; Sequence 12651, Application US/09815242
4      ; Patent No. US20020061569A1
5      ; GENERAL INFORMATION:
6      ; APPLICANT: Haselbeck, Robert
7      ; APPLICANT: Ohlsen, Karl L.
8      ; APPLICANT: Zyskind, Judith W.
9      ; APPLICANT: Wall, Daniel
10     ; APPLICANT: Trawick, John D.
11     ; APPLICANT: Carr, Grant J.
12     ; APPLICANT: Yamamoto, Robert T.
13     ; APPLICANT: Xu, H. Howard
14     ; TITLE OF INVENTION: Identification of Essential Genes in
15     ; FILE OR INVENTION: Prokaryotes
16     ; FILE REFERENCE: ELITRA.01A
17     ; CURRENT APPLICATION NUMBER: US/09/815,242
18     ; CURRENT FILING DATE: 2001-03-21
19     ; PRIOR APPLICATION NUMBER: 60/191,078
20     ; PRIOR FILING DATE: 2000-03-21
21     ; PRIOR APPLICATION NUMBER: 60/206,848
22     ; PRIOR FILING DATE: 2000-05-23
23     ; PRIOR APPLICATION NUMBER: 60/207,727
24     ; PRIOR FILING DATE: 2000-05-26
25     ; PRIOR APPLICATION NUMBER: 60/242,578
26     ; PRIOR FILING DATE: 2000-10-23
27     ; PRIOR APPLICATION NUMBER: 60/253,625
28     ; PRIOR FILING DATE: 2000-11-27
29     ; PRIOR APPLICATION NUMBER: 60/257,931
30     ; PRIOR FILING DATE: 2000-12-22
31     ; PRIOR APPLICATION NUMBER: 60/269,308

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: PRIOR FILING DATE: 2001-02-16
: NUMBER OF SEQ. ID NOS: 14110
: SOFTWARE: FASTSEQ for Windows Version 4.0
: SEQ. ID NO 12651
: LENGTH: 525
: TYPE: PRT
: ORGANISM: Staphylococcus aureus
: US-09-815-242-12651

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Query Match	11.1%	Score 70;	DB 10;	Length 525;
Best Local Similarity	27.9%	Pred. No. 18;		
Matches	29;	Conservative	14;	Mismatches 33; Indels 28; Gaps 5;
QY	35	EAQAQKVDANFYYVILYLMWIGMSEFYVALIVTSVKSRRESDNPFOHYVED-NQ	93	
Db	46	EALAEALYEA-----GRDFRIFYQAPYSRLITLLQMISEPTNYVDSEFS	92	
QY	94	-----EKYSQIL--NLEE--SKATIHENGAAGFMSP	123	
Db	93	VEVEQDENVQKIVYQPLHRIENRNKKLLEAVSRGQIGDGVSP	136	

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RESULT 8
US-10-180-326-1
? Sequence 1, Application US/10180326
? Publication No. US20030049661A1
? GENERAL INFORMATION:
? APPLICANT: Seino, Susumu
? APPLICANT: Shibasaki, Tadao
? APPLICANT: Ozaki, No. US20030049661A1uaki
? TITLE OF INVENTION: Protein Rlm2
? FILE REFERENCE: P21573
? CURRENT APPLICATION NUMBER: US/10/180,326
? CURRENT FILING DATE: 2002-06-27
? PRIOR APPLICATION NUMBER: JP 288372/99
? PRIOR FILING DATE: 1999-10-08<160> 5
? NUMBER OF SEQ ID NOS:
? SOFTWARE: PatentIn version 3.0
? SEQ ID NO 1
? LENGTH: 1590
? TYPE: PRT
? ORGANISM: Mus musculus
? US-10-180-326-1

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Query Match	10.8%	Score 68.5	DB 9	Length 1590
Best Local Similarity	23.9%	Pred. No. 1e-02		
Matches	28	Conservative	19	Mismatches 57; Indels 13; Gaps 3;

  

QY	1	M	S	T	L	E	N	T	O	T	E	V	P	R	I	T	T	D	M	R	O	N	T	A	D	E	A	L	A	V	D	A	N	E	F	Y	L	L	M	A	N	G	F	60
			:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:				
Db	26	M	P	D	L	S	H	T	E	-	E	-	K	I	L	A	M	D	Q	K	E	E	S	V	L	K	E	-	-	-	-	-	-	-	-	-	-	-	-	-	-	72		

  

QY	61	S	P	I	V	A	L	I	V	S	T	A	K	R	R	E	S	N	D	P	Y	Q	I	V	E	D	M	O	E	R	S	Q	L	N	T	E	S	K	A	T	I	H	E	N	G	A	117
		:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:	:						
Db	73	P	F	S	G	I	E	L	V	N	N	N	A	L	P	O	K	O	P	N	E	R	P	O	T	L	H	O	A	F	E	V	A	K	K	G	E	S	O	O	O	E	D	K	G	A	129

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RESULT 9 -
US-09-134-333-12
: Sequence 12, Application US/09134333
: Patent No. US20020076403A1
: GENERAL INFORMATION:
: APPLICANT: LONGACRE-ANDRE, SHIRLEY
: APPLICANT: ROTH, CHARLES
: APPLICANT: NATO, FARIDABANO
: APPLICANT: BARNWELL, JOHN
: APPLICANT: MENDIS, KAMINI
: TITLE OF INVENTION: RECOMBINANT PROTEIN CONTAINING A C-TERMINAL FRAGMENT OF
: TITLE OF INVENTION: PLASMODIUM MSP-1
: FILE REFERENCE: 0660-0135-0XCP
: CURRENT APPLICATION NUMBER: US/09/134,333
: CURRENT FILING DATE: 1999-04-18
: EARLIER APPLICATION NUMBER: PCT/FR97/00290

```

[illegible]

Db 184 H1MESWSPLMNAQILN-DETIKIDAEIG 212

## RESULT 12

US-09-134-333-13  
Sequence 13, Application US/09134333  
Patent No. US20020076403A1

## GENERAL INFORMATION:

APPLICANT: LONGACRE-ANDRE, SHIRLEY

APPLICANT: ROTH, CHARLES

APPLICANT: NATO, FARIDABANO

APPLICANT: BARWELL, JOHN

APPLICANT: MENDIS, KAMINI

TITLE OF INVENTION: RECOMBINANT PROTEIN CONTAINING A C-TERMINAL FRAGMENT OF

TITLE OF INVENTION: PLASMODIUM MSP-1

FILE REFERENCE: 0660-0135-0KCIIP

CURRENT APPLICATION NUMBER: US/09/134,333

CURRENT FILING DATE: 1999-04-18

EARLIER APPLICATION NUMBER: PCT/FR97/00290

EARLIER FILING DATE: 1997-02-14

EARLIER APPLICATION NUMBER: FR96/01822

EARLIER FILING DATE: 1996-02-14

NUMBER OF SEQ ID NOS: 14

SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 13

LENGTH: 380

TYPE: PRT

ORGANISM: Plasmodium vivax-like sp.

FEATURE:

OTHER INFORMATION: Amino Acids 1-140-REGION I

FEATURE:

OTHER INFORMATION: Amino Acids 141-178-REGION II

FEATURE:

OTHER INFORMATION: Amino Acids 179-283-REGION III

FEATURE:

OTHER INFORMATION: Amino Acids 284-380-REGION IV

US-09-134-333-13

Query Match 10.5%; Score 66.5; DB 10; Length 380;

Best Local Similarity 25.2%; Pred. No. 29;

Matches 32; Conservative 18; Mismatches 26; Indels 51; Gaps 7;

QY 28 QNTT--AEQALQAKVAENFYVILYLMWIGMFSLIYALIVSYKSKRRENSN--- 81

Db 2 QVTTGEASEAPELVIPAGISDYVYTKPLAGAT-----KTIKKOLEHNVAEMT 52

QY 82 -----DPYH-----QYVEDMQEKYKSQLNLESKATI-- 110

Db 53 NITDMLDSRLKKRNYFLEVLSNDLNPFRYSSSGEYIIKD--PYK--LIDLEKRLIGS 107

QY 111 HENIGAA 117

Db 108 KYKIGAS 114

## RESULT 13

US-09-815-242-5778

Sequence 5778, Application US/09815242

Patent No. US20020061569A1

## GENERAL INFORMATION:

APPLICANT: Haselbeck, Robert

APPLICANT: Ohlsen, Karl L.

APPLICANT: Zyskind, Judith W.

APPLICANT: Wall, Daniel

APPLICANT: Trawick, John D.

APPLICANT: Carr, Grant J.

APPLICANT: Yamamoto, Robert T.

TITLE OF INVENTION: Identification of Essential Genes in

TITLE OF INVENTION: Prokaryotes

FILE REFERENCE: ELITRA.011A

CURRENT APPLICATION NUMBER: US/09/815,242

CURRENT FILING DATE: 2001-03-21  
PRIOR APPLICATION NUMBER: 60/191,078  
PRIOR FILING DATE: 2000-03-21  
PRIOR APPLICATION NUMBER: 60/206,848  
PRIOR FILING DATE: 2000-05-23  
PRIOR APPLICATION NUMBER: 60/207,727  
PRIOR FILING DATE: 2000-05-26  
PRIOR APPLICATION NUMBER: 60/242,578  
PRIOR FILING DATE: 2000-10-23  
PRIOR APPLICATION NUMBER: 60/253,625  
PRIOR FILING DATE: 2000-11-27  
PRIOR APPLICATION NUMBER: 60/257,931  
PRIOR FILING DATE: 2000-12-22  
PRIOR APPLICATION NUMBER: 60/269,308  
PRIOR FILING DATE: 2001-02-16  
NUMBER OF SEQ ID NOS: 14110  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 5778

LENGTH: 553

TYPE: PRT

ORGANISM: Staphylococcus aureus

US-09-815-242-5778

Query Match 10.4%; Score 65.5; DB 10; Length 553;

Best Local Similarity 20.8%; Pred. No. 39;

Matches 26; Conservative 24; Mismatches 44; Indels 31; Gaps 5;

QY 9 QTEDEVFRRIT---FTYNDNRONTTAEQALQAKVAENFYVILY----- 52

Db 190 QDINDVTEAKTKNSFPVLQAGMSSSAEETNAIKLYERNLNPFVETFGAGYSKELN 249

QY 53 -LAWMIGMF-----SEIYALIVSYKSKRRENSDPYHQYVEDMQEKYKSQLNLE 105

Db 250 HFGRGVGLFRNQYGDLELRSDLVATI-----GDPIL-EYASNMNKELETOIINDE 301

QY 106 SKATI 110

Db 302 VQAEI 306

## RESULT 14

US-09-982-610-24

Sequence 24, Application US/09982610

Patent No. US20020146420A1

## GENERAL INFORMATION:

APPLICANT: Genentech, Inc.

APPLICANT: Bennett, Brian D.

APPLICANT: Goeddel, David

APPLICANT: Lee, James M.

APPLICANT: Matthews, William

APPLICANT: Tsai, Siao Ping

APPLICANT: Wood, William I.

TITLE OF INVENTION: PROTEIN TYROSINE KINASE AGONIST ANTIBODIES

NUMBER OF SEQUENCES: 45

CORRESPONDENCE ADDRESS:

ADDRESSEE: Genentech, Inc.

STREET: 460 Point San Bruno Blvd

CITY: South San Francisco

STATE: California

COUNTRY: USA

ZIP: 94080

COMPUTER READABLE FORM:

MEDIUM TYPE: 3.5 Inch, 1.44 Mb floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Winpatin (Genentech)

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/982,610

FILING DATE: 17-Oct-2001

CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/446,648

FILING DATE: 1996-MAY-23



```

; APPLICATION NUMBER: 08/222616
; FILING DATE: 04-APR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Lee, Wendy M.
; REGISTRATION NUMBER: 40,378
; REFERENCE/DOCKET NUMBER: P0821P3PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415/225-1994
; TELEFAX: 415/952-9881
; TELE: 910/371-7168
; INFORMATION FOR SEQ ID NO: 24:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1276 amino acids
; TYPE: Amino Acid
; TOPOLOGY: Linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 24:
US-09-982-610-24

Query Match      10.4%; Score 65.5; DB 10; Length 1276;
Best Local Similarity 28.2%; Pred. No. 1.6e+02;
Matches 31; Conservative 13; Mismatches 37; Indels 29; Gaps 5;

QY 24 DNMKONTAEQALQAKDAENFYVILYLMWIGMFSTIYVAILVSTYKRRR-HSND 82
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
DB 532 EGMK-----EQALTAGTAVG---VVLVLY-----IVYAVLCLRKOSNGRAEYSD 576
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
QY 83 PYHOYIV-----EDMOEKYSQILNLESKATIHENIGAGE 119
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
DB 577 KKGQYILGIGTKYIDPTEYDPAVNEAFNAKEIDVSTYKIEVIGAGEF 626

RESULT 15
US-09-896-994-2
; Sequence 2, Application US/09896994
; Publication No. US20030074024A1
; GENERAL INFORMATION:
; APPLICANT: Ken Stokes
; Jos e Morissette
; TITLE OF INVENTION: SYSTEMS AND METHODS FOR ENHANCING CARDIAC
; SIGNAL SENSING BY CARDIAC PACEMAKERS THROUGH GENETIC TREATM
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Woodcock Washburn Kurtz Mackiewicz and No. US20030074024A1ris
; STREET: One Liberty Place - 46th Floor
; CITY: Philadelphia
; STATE: PA
; COUNTRY: U.S.A.
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WordPerfect 6.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/896,994
; FILING DATE: 02-Jul-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/514,907
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul K. Legaard
; REGISTRATION NUMBER: 38,534
; REFERENCE/DOCKET NUMBER: MEDT-0029/P-3586
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-3100
; TELEFAX: (215) 568-3439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2016 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: unknown

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; SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-09-896-994-2

Query Match      10.4%; Score 65.5; DB 9; Length 2016;
Best Local Similarity 20.5%; Pred. No. 2.9e+02;
Matches 26; Conservative 27; Mismatches 43; Indels 31; Gaps 6;

QY 4 LSNFTQLEDVFRIRFIYMDNMKONTAEQALQAKDAENFYVILYLMWIGMFSTI 63
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
DB 363 LALFRLMTQDCWKERY-----QQTLSR---AKITMIFPMVIFLG--SFY 403
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
QY 64 IVALIVSTYKSKRRRHSNDPYHOYIVEDMOEKYK-----SQILNLESKATIH--ENIGA 116
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
DB 404 LVNLIYAVVAMAYEEQ-----NQATIAETEKEKRFQEMELKKHEHALTRGVDTYSR 458
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
QY 117 AGFKMSP 123
   :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :  :
DB 459 SLEWMS 465

Search completed: May 15, 2003, 14:35:09
Job time : 73 secs

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XX Novel nucleic acids encoding M1RP1, M1RP2 and M1RP3, useful for
PT diagnosing and treating ion channel disorders, especially long QT
PR syndrome.
XX
XX Claim 19; Page 119; 132pp; English.
XX
CC The invention relates to novel ion channel proteins related to
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of
CC the invention are human and rat KCNE2 (M1RP1; AAB29585 and AAB29586,
CC respectively); human and mouse KCNE3 (M1RP2; AAB29587 and AAB29588,
CC respectively); human and mouse KCNE4 (M1RP3; AAB29589 and AAB29590,
CC respectively). The CDNA encoding these proteins are given in AAC64071-
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier
CC potassium channels (I-KR), mutations in which are associated with long
CC QT syndrome. The invention also relates to methods of diagnosing long QT
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic
CC nonhuman animals comprising a heterologous ion channel protein gene
CC of the invention, a transgenic animal comprising human KCNE2 and HERG
CC DNA, and methods of and screening drugs for treating long QT syndrome
CC using KCNE2 proteins (including mutants), nucleic acids encoding them
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic
CC acids, and proteins may be used for diagnosing or treating ion channel
CC disorders, especially long QT syndrome. Transgenic animals comprising
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.
CC The present sequence represents human KCNE2 (M1RP1).
XX
SQ Sequence 123 AA:
Query Match 100.0%; Score 632; DB 21; Length 123;
Best Local Similarity 100.0%; Pred. No. 1.9e-67;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 MSTLSNFTQLEDFERRITFYMDNMRONTAEOALQAKVDAENFYVILYLMWIGMF 60
DB 1 MSTLSNFTQLEDFERRITFYMDNMRONTAEOALQAKVDAENFYVILYLMWIGMF 60
OY 61 SFTIYALIVSVTKSKRREHSNDPYHQYIYEDMQEKYSQILNLESKATIHENIGAGFK 120
DB 61 SFTIYALIVSVTKSKRREHSNDPYHQYIYEDMQEKYSQILNLESKATIHENIGAGFK 120
OY 121 MSP 123
DB 121 MSP 123
XX
RESULT 2
AAM78528
ID AAM78528 standard; Protein: 123 AA.
XX
AC AAM78528;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human protein SEQ ID NO 1190.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation.
XX
OS Homo sapiens.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US04098.
XX
PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.

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PR 19-JUN-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QA, Wang D, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejrtman T, Goodrich R.
XX
DR WPI: 2001-476283/51.
DR N-PSDB; AAK51661.
XX
PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
XX
PS Claim 20; Page 3438-3439; 6221pp; English.
XX
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666
CC (AAM80020) are omitted as the relevant pages from the sequence listing
CC were missing at the time of publication.
XX
SQ Sequence 123 AA:
Query Match 100.0%; Score 632; DB 22; Length 123;
Best Local Similarity 100.0%; Pred. No. 1.9e-67;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 MSTLSNFTQLEDFERRITFYMDNMRONTAEOALQAKVDAENFYVILYLMWIGMF 60
DB 1 MSTLSNFTQLEDFERRITFYMDNMRONTAEOALQAKVDAENFYVILYLMWIGMF 60
OY 61 SFTIYALIVSVTKSKRREHSNDPYHQYIYEDMQEKYSQILNLESKATIHENIGAGFK 120
DB 61 SFTIYALIVSVTKSKRREHSNDPYHQYIYEDMQEKYSQILNLESKATIHENIGAGFK 120
OY 121 MSP 123
DB 121 MSP 123
XX
RESULT 3
AAM18627
ID AAM18627 standard; Protein: 123 AA.
XX
AC AAM18627;
XX
DT 12-OCT-2001 (first entry)
XX
DE Peptide #5061 encoded by probe for measuring cervical gene expression.
XX
KW Probe; human; microarray; gene expression; cervical epithelial cell;
KW cervical cancer.
XX
OS Homo sapiens.
XX
PN WO200157278-A2.
XX
PD 09-AUG-2001.
XX

```

PF 30-JAN-2001; 2001WO-US006070.  
XX  
PR 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX WPI; 2001-488901/53.  
XX  
DR WPI; 2001-488901/53.  
XX  
PT Human genome-derived single exon nucleic acid probes useful for  
PS analyzing gene expression in human cervical epithelial cells -  
XX  
PS Claim 27; SEQ ID No 23453; 487bp; English.  
XX  
CC The present invention relates to human single exon nucleic acid probes  
CC (SENP; see A110068-A1128459). The present sequence is a peptide encoded  
CC by one such probe. The SENPs are derived from human Hela cells. The SENPs  
CC can be used to produce a single exon microarray, which can be used for  
CC measuring human gene expression in a sample derived from human cervical  
CC epithelial cells. By measuring gene expression, the probes are therefore  
CC useful in grading and/or staging of diseases of the cervix, notably  
CC cervical cancer.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 123 AA;

Query Match 100.0%; Score 632; DB 22; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQTEDEYFRRIFFITTYMDNWRONTAEQALQAKYDAENFYVILYLMVMIGMF 60  
DB 1 MSTLSNFTQTEDEYFRRIFFITTYMDNWRONTAEQALQAKYDAENFYVILYLMVMIGMF 60  
QY 61 SFTIYAILVSTYVSKRRHSNDPYHQYIYEDWQEKYSQILNLEESKATIHENIGAGGFK 120  
DB 61 SFTIYAILVSTYVSKRRHSNDPYHQYIYEDWQEKYSQILNLEESKATIHENIGAGGFK 120  
QY 121 MSP 123  
DB 121 MSP 123

RESULT 4  
AAM06194  
ID AAM06194 standard; Protein; 123 AA.  
XX  
AC AAM06194;  
XX  
DT 09-OCT-2001 (first entry)  
XX  
DE Peptide #4876 encoded by probe for measuring breast gene expression.  
XX  
KW Probe: human; breast disease; breast cancer; development disorder;  
KW inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
XX  
OS Homo sapiens.  
XX  
PN WO200157270-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 29-JAN-2001; 2001WO-US00661.  
XX

PR 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX WPI; 2001-476286/51.  
XX  
DR WPI; 2001-476286/51.  
XX  
PT Novel single exon nucleic acid probe used to measuring gene expression  
PS in a human breast -  
XX  
PS Claim 27; SEQ ID No 14934; 322bp; English.  
XX  
CC The present invention relates to novel single exon nucleic acid probes  
CC (see A1100010-A110067). The present sequence is a peptide encoded by one  
CC such probe. The probes are useful for measuring human gene expression in  
CC a human breast sample, where the probe hybridises at high stringency to a  
CC nucleic acid expressed in the human breast. The probes are useful for  
CC predicting, diagnosing, grading, staging, monitoring and prognosing  
CC diseases of the human breast, particularly those diseases with polygenic  
CC aetiology. The diseases include: breast cancer, disorders of development,  
CC inflammatory diseases of the breast, fibrocystic changes, proliferative  
CC breast disease and non-carcinoma tumours.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 123 AA;

Query Match 100.0%; Score 632; DB 22; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQTEDEYFRRIFFITTYMDNWRONTAEQALQAKYDAENFYVILYLMVMIGMF 60  
DB 1 MSTLSNFTQTEDEYFRRIFFITTYMDNWRONTAEQALQAKYDAENFYVILYLMVMIGMF 60  
QY 61 SFTIYAILVSTYVSKRRHSNDPYHQYIYEDWQEKYSQILNLEESKATIHENIGAGGFK 120  
DB 61 SFTIYAILVSTYVSKRRHSNDPYHQYIYEDWQEKYSQILNLEESKATIHENIGAGGFK 120  
QY 121 MSP 123  
DB 121 MSP 123

RESULT 5  
AAB67800  
ID AAB67800 standard; Protein; 123 AA.  
XX  
AC AAB67800;  
XX  
DT 29-JUN-2001 (first entry)  
XX  
DE Amino acid sequence of human potassium channel subunit Isk2.  
XX  
KW Human; potassium channel; Isk2; gene therapy; gastric motility;  
KW gastric acid secretion; anti-arrhythmic agent; myocardial infarction.  
XX  
OS Homo sapiens.  
XX  
PN WO200127246-A1.  
XX  
PD 19-APR-2001.  
XX  
PF 10-OCT-2000; 2000WO-US28014.  
XX

PR 12-OCT-1999; 99US-0158781.  
XX  
PA (MERI ) MERCK & CO INC.  
XX  
PI Swanson RJ, Liu Y, Folander K;  
XX WPI; 2001-273764/28.  
DR N-PSDB; AAF80269.  
XX  
PT New DNA encoding the Isk2 potassium channel subunit, useful e.g. for  
XX detecting mutations and screening for therapeutic agents -  
PS Claim 8; Fig 1b; 46pp; English.  
XX  
XX The present sequence represents a human potassium channel subunit,  
CC designated Isk2. The Isk2 polynucleotide, and derived probes, are  
CC designed to detect mutations in the Isk2 gene, to determine  
CC levels of mRNA expression and to isolate homologous sequences; for  
CC recombinant expression of Isk2; in gene therapy to increase potassium  
CC channel activity and to generate transgenic animals, as models and  
CC for drug screening. Recombinant Isk2 is used for studying biochemical  
CC activity of Isk2 and its role in disorders of gastric motility and  
CC gastric acid secretion, and to raise specific antibodies. Isk2  
CC modulators are potentially useful for treating diseases associated with  
CC increased or reduced potassium channel activity e.g. as  
CC anti-arrhythmic agents for treating myocardial infarction and as  
CC regulators of gastric acid secretion.  
XX  
SQ Sequence 123 AA;  
XX  
Query Match 100.0%; Score 632; DB 22; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 MSTLSNFTQLEDFVFRIFITTYMDNMRONTTAEQALQAKVDENFYVILYLMWIGMF 60  
DB 1 MSTLSNFTQLEDFVFRIFITTYMDNMRONTTAEQALQAKVDENFYVILYLMWIGMF 60  
OY 61 SFTIIVAILVSTYKSKRREHSNDPIHOYIVEDMQEKYSQILNLESKATIHENIGAAGFK 120  
DB 61 SFTIIVAILVSTYKSKRREHSNDPIHOYIVEDMQEKYSQILNLESKATIHENIGAAGFK 120  
OY 121 MSP 123  
DB 121 MSP 123  
XX  
RESULT 6  
AAU00215  
ID AAU00215 standard; Protein; 123 AA.  
XX  
AC AAU00215;  
XX  
DT 10-MAY-2001 (first entry)  
XX  
DE Human potassium channel regulatory protein, MINK2.  
XX  
KW Human; MINK2; potassium channel; cardiac arrhythmia; hypertension;  
KW angina; asthma; diabetes; renal insufficiency; urinary incontinence;  
KW irritable colon; epilepsy; cerebrovascular ischemia; autoimmune disease.  
XX  
OS Homo sapiens.  
XX  
PN WO200114403-A1.  
XX  
PD 01-MAR-2001.  
XX  
PF 18-AUG-2000; 2000MO-US22799.  
XX  
PR 20-AUG-1999; 99US-0379201.  
XX  
PA (UYCA-) UNIV CASE WESTERN RESERVE.  
XX

PI Flicker E, Mible B, Brown AM;  
XX  
DR WPI; 2001-218424/22.  
DR N-PSDB; AAS00245.  
XX  
PT Novel potassium channel gene termed Mink2 encoding potassium channel  
XX regulatory protein, useful for screening compounds that are useful for  
XX treating diseases caused by aberrant potassium activity -  
PS Disclosure; Fig 9; 39pp; English.  
XX  
XX The sequence represents the amino acid sequence of human potassium  
CC channel regulatory protein, MINK2. MINK2 sequence is useful for producing  
CC a potassium channel regulatory protein useful for in vitro or in vivo  
CC screening of agonistic or antagonistic compounds that are useful for  
CC treating diseases caused by aberrant potassium activity, such as human  
CC cardiac arrhythmias, hypertension, angina, asthma, diabetes, renal  
CC insufficiency, urinary incontinence, irritable colon, epilepsy,  
CC cerebrovascular ischemia, and autoimmune disease.  
XX  
SQ Sequence 123 AA;  
XX  
Query Match 100.0%; Score 632; DB 22; Length 123;  
Best Local Similarity 100.0%; Pred. No. 1.9e-67;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 MSTLSNFTQLEDFVFRIFITTYMDNMRONTTAEQALQAKVDENFYVILYLMWIGMF 60  
DB 1 MSTLSNFTQLEDFVFRIFITTYMDNMRONTTAEQALQAKVDENFYVILYLMWIGMF 60  
OY 61 SFTIIVAILVSTYKSKRREHSNDPIHOYIVEDMQEKYSQILNLESKATIHENIGAAGFK 120  
DB 61 SFTIIVAILVSTYKSKRREHSNDPIHOYIVEDMQEKYSQILNLESKATIHENIGAAGFK 120  
OY 121 MSP 123  
DB 121 MSP 123  
XX  
RESULT 7  
AAU99168  
ID AAU99168 standard; Protein; 123 AA.  
XX  
AC AAU99168;  
XX  
DT 24-SEP-2002 (first entry)  
XX  
DE Human ether-a-go-go related interacting protein M1RP1.  
XX  
KW Human; human ether-a-go-go related gene; HERG; KCR1; M1RP1;  
KW long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;  
KW potassium channel.  
XX  
OS Homo sapiens.  
XX  
PN WO200242735-A2.  
XX  
PD 30-MAY-2002.  
XX  
PF 30-OCT-2001; 2001MO-US45644.  
XX  
PR 30-OCT-2000; 2000US-244340P.  
XX  
PA (UYVA-) UNIV VANDERBILT.  
XX  
PI Balser JR, George AL, Roden DM;  
XX  
DR WPI; 2002-527650/56.  
XX  
DR N-PSDB; ABR86573.  
XX  
PT Identifying a potassium channel activity modulator for drug design,  
XX comprises contacting a compound with a potassium channel and rat  
XX cerebellar cDNA library (KCR1) polypeptide, and determining activity -

XX Disclousre: Page 163; 164pp; English.

XX The invention relates to identifying (M1) a compound that modulates  
 CC biological activity of a potassium channel (PC), by contacting a polypeptide  
 CC compound with a structure comprising a PC polypeptide and a polypeptide  
 CC cloned from a rat cerebellar cDNA library (KCRI) and determining the  
 CC activity of the PC polypeptide in the presence and absence of the  
 CC compound, where a difference in the activities indicates modulation of  
 CC biological activity of PC. Also include are identifying (M2) a candidate  
 CC compound that modulates the biological activity of a complex comprising a  
 CC human ether-a-go-go-related gene (HERG) channel polypeptide and a KCRI  
 CC polypeptide, identifying (M3) a candidate compound as a modulator of KCRI  
 CC expression, modulating (M4) PC function in a subject, comprising  
 CC administering to the subject a substance that provides expression of a  
 CC KCRI-encoding nucleic acid molecule in a cell or tissue, where modulated  
 CC PC function is desired, screening (M5) for susceptibility to a drug-  
 CC induced cardiac arrhythmia in a subject, comprising obtaining a  
 CC biological sample from the subject and detecting a polymorphism of a KCRI  
 CC gene in the biological sample from the subject, where the presence of the  
 CC polymorphism indicates the susceptibility of the subject to a  
 CC drug-induced cardiac arrhythmia, an oligonucleotide pair, where a first  
 CC oligonucleotide of the pair hybridises to a first portion of a KCRI gene  
 CC which includes a polymorphism of the KCRI gene, and the second  
 CC oligonucleotide of the pair hybridises to a second portion of the KCRI  
 CC gene that is adjacent to the first portion and a set of antisense  
 CC oligonucleotide primers, suitable for amplifying a portion of a KCRI gene  
 CC which includes a polymorphism of the KCRI gene, (M1) is useful for  
 CC identifying a compound that modulates biological activity of PC,  
 CC especially HERG, for modulating PC function (i.e. modulating HERG  
 CC activity) in a mammal, by preparing a composition comprising the  
 CC compound and administering the composition. The compound is useful for  
 CC treating or preventing long QT syndrome (LQTS) and is useful in drug  
 CC designing. The present sequence represents a HERG interacting  
 CC protein MiRP1 (not defined).

XX Sequence 123 AA;

Query Match 100.0%; Score 632; DB 23; Length 123;

Best Local Similarity 100.0%; Pred. No. 1.9e-67; Mismatches 0; Indels 0; Gaps 0;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQTLDEYFRRIFFITTYMDNMWRONTTAEQALQAKYDAENFYVILLYLMVMIGMF 60  
 DB 1 MSTLSNFTQTLDEYFRRIFFITTYMDNMWRONTTAEQALQAKYDAENFYVILLYLMVMIGMF 60

QY 61 SFTIYAILVSTYKSKRRHSNDPYHQYIYEDMOKERYKSOILNLEESKATIHENIGAAGFK 120  
 DB 61 SFTIYAILVSTYKSKRRHSNDPYHQYIYEDMOKERYKSOILNLEESKATIHENIGAAGFK 120

QY 121 MSP 123  
 DB 121 MSP 123

RESULT 8

AAE22095

ID AAE22095 standard; Protein; 123 AA.

XX AAE22095;

XX 25-JUL-2002 (first entry)

XX Human MiRP1 wild type protein.

XX Human; Min-K related ion channel protein; MiRP1; ion channel disorder;

XX KCNE2; long QT syndrome; LQTS; cardiac arrhythmia.

OS Homo sapiens.

XX WO200222875-A2.

XX 21-MAR-2002.

XX 11-SEP-2001; 2001MO-US28332.

XX 11-SEP-2000; 2000US-231571P.

XX (UYVA ) UNIV YALE.

XX Goldstein SAN;

XX WPI; 2002-362360/39.

XX N-PSDB; AAD35170.

XX Novel gene encoding Min-K related ion channel protein subunit and

XX polymorphisms in this gene associated with antibiotic-induced long QT

XX syndrome -

XX Claim 8; Fig 1A; 49pp; English.

XX The present invention relates to novel KCNE2 genes encoding Min-K related  
 CC (MiRP) 1 ion channel proteins and polymorphisms in these genes that are  
 CC associated with ion channel disorders including antibiotic-induced long  
 CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
 CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCNE2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human MiRP1 wild type protein.

XX Sequence 123 AA;

Query Match 100.0%; Score 632; DB 23; Length 123;

Best Local Similarity 100.0%; Pred. No. 1.9e-67; Mismatches 0; Indels 0; Gaps 0;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQTLDEYFRRIFFITTYMDNMWRONTTAEQALQAKYDAENFYVILLYLMVMIGMF 60  
 DB 1 MSTLSNFTQTLDEYFRRIFFITTYMDNMWRONTTAEQALQAKYDAENFYVILLYLMVMIGMF 60

QY 61 SFTIYAILVSTYKSKRRHSNDPYHQYIYEDMOKERYKSOILNLEESKATIHENIGAAGFK 120  
 DB 61 SFTIYAILVSTYKSKRRHSNDPYHQYIYEDMOKERYKSOILNLEESKATIHENIGAAGFK 120

QY 121 MSP 123  
 DB 121 MSP 123

RESULT 9

ABB11948

ID ABB11948 standard; peptide; 135 AA.

XX ABB11948;

XX 11-JAN-2002 (first entry)

XX Human MiRP1 homologue, SEQ ID NO:2318.

XX Human; cytokine; cell proliferation; cell differentiation; growth factor;

XX haematopoiesis regulation; tissue growth; immunomodulator; activin;

XX inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;

XX proliferation; metastasis; cancer; tumour; haematopoietic disorder;

XX myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;

XX chronic inflammatory condition; proliferative retinopathy;

XX atherosclerosis; coronary heart disease; arterial ischemia;

XX bone disorder; osteoporosis; vascular growth disorder;

XX tissue regeneration; wound healing; infection; immune disorder;

XX cell culture; drug screening; gene therapy; anti-inflammatory;

XX antiasthmatic; antiarthritis; haemostatic; antiarteriosclerotic;

XX cytoskeletal; osteopathic; vasotropic; cardiac; virucide; antibacterial;

XX antifungal; vulnery; antidiabetic.

XX Homo sapiens.

XX MO200157188-A2.  
 PN 09-AUG-2001.  
 PD 09-AUG-2001.  
 XX 05-FEB-2001; 2001WO-US03800.  
 PF 03-FEB-2000; 2000US-0496914.  
 XX 27-APR-2000; 2000US-0560875.  
 XX (HYSE-) HYSEQ INC.  
 PA Tang YT, Liu C, Drmanac RT;  
 PI WPI: 2001-457740/49.  
 DR N-PSDB; ABA09192.  
 XX  
 XX Human proteins and DNA encoding sequences useful for preventing,  
 PT treating or ameliorating a medical condition in a mammalian subject  
 PT e.g. arthritis and cancer -  
 XX  
 PS Claim 20; Page 283; 1963pp; English.  
 XX  
 XX Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and  
 CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The  
 CC invention also relates to vectors and recombinant host cells comprising a  
 CC nucleotide of the invention, methods of producing the novel polypeptides,  
 CC antibodies against the polypeptides, methods of detecting the nucleotides  
 CC or polypeptides in a sample, and methods of identifying compounds which  
 CC bind to polypeptides of the invention. Although novel, many of the  
 CC polypeptides of the invention have homology to known proteins, thereby  
 CC giving an insight into their probable biological activities, and hence  
 CC potential therapeutic applications. The polypeptides of the invention may  
 CC have various activities, including cytokine, cell proliferation or cell  
 CC differentiation activities; stem cell growth factor activity;  
 CC hematopoietic regulatory activity; tissue growth activity;  
 CC immunomodulatory activity; activin or inhibin-related activities;  
 CC chemotactic or chemokinetic activities; haemostatic, thrombotic or  
 CC thrombolytic activities; receptor or ligand activities; or may be  
 CC involved in oncogenesis, cancer cell proliferation or metastasis.  
 CC Depending on their biological activities, polypeptides and nucleotides of  
 CC the invention are useful for preventing, treating or ameliorating medical  
 CC conditions, e.g., by protein or gene therapy. Such conditions include  
 CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell  
 CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),  
 CC proliferative retinopathy, atherosclerosis, coronary heart disease,  
 CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal  
 CC vascular growth. Polypeptides involved with tissue regeneration and  
 CC repair (or nucleic acids encoding them) may be used to promote wound  
 CC healing (e.g., of burns, incisions and ulcers), while those with  
 CC immunomodulatory activities may be used in the treatment of viral,  
 CC bacterial and fungal infections in addition to immune disorders.  
 CC Polypeptides with growth factor activity may be used in cell cultures to  
 CC promote cell growth. For example, such polypeptides may be used to  
 CC manipulate stem cells in culture to give rise to neuroepithelial cells  
 CC that can be used to augment or replace cells damaged by illness,  
 CC autoimmune disease or accidental damage. The polypeptides and nucleotides  
 CC may also be used in the diagnosis of the above conditions, and in drug  
 CC screening techniques. The present sequence represents a novel human  
 CC polypeptide of the invention.  
 XX  
 SQ Sequence 135 AA;  
 Query Match 100.0%; Score 632; DB 22; Length 135;  
 Best Local Similarity 100.0%; Pred. No. 2,2e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 MSTLSNFTQLEDFVFRRIFFTYMDNMRQNTTADQALQAKVDAENFYVILYLVAMWIGMF 60  
 DB 13 MSTLSNFTQLEDFVFRRIFFTYMDNMRQNTTADQALQAKVDAENFYVILYLVAMWIGMF 72  
 QY 61 SFTIIVALTSTVSKRRHSNDPYHQYIVEDWQEKYSQILMLESKATIHENIGAGPK 120  
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||

DB 73 SFTIIVALTSTVSKRRHSNDPYHQYIVEDWQEKYSQILMLESKATIHENIGAGPK 132  
 QY 121 MSP 123  
 ||||  
 DB 133 MSP 135  
 RESULT 10  
 ID AAM79512  
 ID AAM79512 standard; Protein; 135 AA.  
 XX  
 XX AAM79512;  
 XX  
 XX 06-NOV-2001 (first entry)  
 XX  
 XX Human protein SEQ ID NO 3158.  
 DE  
 XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;  
 KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;  
 KW tissue growth factor; immunomodulatory; cancer; leukaemia;  
 KW nervous system disorder; arthritis; inflammation.  
 XX  
 OS Homo sapiens.  
 XX  
 XX WO200157190-A2.  
 PN 09-AUG-2001.  
 PD 09-AUG-2001.  
 XX  
 PF 05-FEB-2001; 2001WO-US04098.  
 XX  
 XX 03-FEB-2000; 2000US-0496914.  
 PR 27-APR-2000; 2000US-0560875.  
 PR 20-JUN-2000; 2000US-0598075.  
 PR 19-JUL-2000; 2000US-0620325.  
 PR 01-SEP-2000; 2000US-0654936.  
 PR 15-SEP-2000; 2000US-0663561.  
 PR 20-OCT-2000; 2000US-0693325.  
 PR 30-NOV-2000; 2000US-0728422.  
 XX  
 PA (HYSE-) HYSEQ INC.  
 XX  
 PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
 PI Zhao QH, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;  
 PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;  
 DR WPI: 2001-476283/51.  
 XX N-PSDB; AAK52645.  
 PT Nucleic acids encoding polypeptides with cytokine-like activities,  
 PT useful in diagnosis and gene therapy -  
 XX  
 PS Claim 20; Page 270; 6221pp; English.  
 XX  
 XX The invention relates to polynucleotides (AAK51456-AAK53435) and the  
 CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to  
 CC cytokine, cell proliferation or cell differentiation or which may induce  
 CC production of other cytokines in other cell populations. The  
 CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
 CC peptide therapy. The polypeptides have various cytokine-like activities,  
 CC e.g. stem cell growth factor activity, haematopoiesis regulating  
 CC activity, tissue growth factor activity, immunomodulatory activity and  
 CC activin/inhibin activity and may be useful in the diagnosis and/or  
 CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
 CC inflammation.  
 CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
 CC (AAK80020) are omitted as the relevant pages from the sequence listing  
 CC were missing at the time of publication.  
 XX  
 SQ Sequence 135 AA;  
 Query Match 100.0%; Score 632; DB 22; Length 135;  
 Best Local Similarity 100.0%; Pred. No. 2,2e-67;  
 Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;



QY 1 MSTLSNFTQTLIEDVFRIRIFTYMDNMKONTTAEQALQAKVDAENFYVILYLMVMGMF 60  
 DB 13 MSTLSNFTQTLIEDVFRIRIFTYMDNMKONTTAEQALQAKVDAENFYVILYLMVMGMF 72  
 QY 61 SFIIVAILVSTYVSKRRRHSNDPYHQYIVEDMQEYKSQLNLEESKATIHENIGAAGFK 120  
 DB 73 SFIIVAILVSTYVSKRRRHSNDPYHQYIVEDMQEYKSQLNLEESKATIHENIGAAGFK 132  
 QY 121 MSP 123  
 DB 133 MSP 135

RESULT 11  
 AAB29593  
 ID AAB29593 standard; Protein; 123 AA.  
 XX AAB29593;  
 AC  
 XX  
 DT 19-FEB-2001 (first entry)  
 XX  
 DE Human potassium channel protein KCNE2 (MIRP1) mutant Q9E.  
 XX  
 KW Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
 KW MinK-related; long QT syndrome; cardiac arrhythmia;  
 KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KW HERG; mutant; muteln.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 PN MO20006343-A1.  
 XX  
 PD 26-OCT-2000.  
 XX  
 PF 14-APR-2000; 2000WO-US10004.  
 XX  
 PR 15-APR-1999; 99US-0129404.  
 XX  
 PA (UTAH ) UNIV UTAH RES FOUND.  
 PA (UYVA ) UNIV YALE.  
 XX  
 PI Abbot GW, Sestl F, Splawski I, Keating MT, Goldstein SAN;  
 PI WPI: 2000-672747/65.  
 DR N-PSDB; AAC64083.  
 XX  
 PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 XX  
 PS Claim 56; Page -; 132pp; English.  
 XX  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (MinK) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MIRP1; AAB29586 and AAB29588,  
 CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-Kr), mutations in which are associated with long QT  
 CC syndrome. The invention, also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising

CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents a mutant human KCNE2 (MIRP1)  
 CC specifically claimed for use in diagnostic and drug screening methods of  
 CC the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 protein sequence shown on page  
 CC 119.  
 XX  
 SQ Sequence 123 AA;  
 Query Match 99.5%; Score 629; DB 21; Length 123;  
 Best Local Similarity 99.2%; Pred. No. 4,4e-67;  
 Matches 122; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MSTLSNFTQTLIEDVFRIRIFTYMDNMKONTTAEQALQAKVDAENFYVILYLMVMGMF 60  
 DB 1 MSTLSNFTQTLIEDVFRIRIFTYMDNMKONTTAEQALQAKVDAENFYVILYLMVMGMF 60  
 QY 61 SFIIVAILVSTYVSKRRRHSNDPYHQYIVEDMQEYKSQLNLEESKATIHENIGAAGFK 120  
 DB 61 SFIIVAILVSTYVSKRRRHSNDPYHQYIVEDMQEYKSQLNLEESKATIHENIGAAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123

RESULT 12  
 AAE22094  
 ID AAE22094 standard; Protein; 123 AA.  
 XX AAE22094;  
 AC  
 XX  
 DT 25-JUL-2002 (first entry)  
 XX  
 DE Human MIRP1 mutant protein (A116V).  
 XX  
 KW Human; Min-K related ion channel protein; MIRP1; ion channel disorder;  
 KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; muteln.  
 XX  
 OS Homo sapiens.  
 OS  
 FH Key Location/Qualifiers  
 FT Misc-difference 116  
 FT /note= "Wild type Ala substituted with Val"  
 XX  
 PN WO200222875-A2.  
 XX  
 PD 21-MAR-2002.  
 XX  
 PF 11-SEP-2001; 2001WO-US28332.  
 XX  
 PR 11-SEP-2000; 2000US-231571P.  
 XX  
 PA (UYVA ) UNIV YALE.  
 PA Goldstein SAN;  
 PI WPI: 2002-362360/39.  
 DR N-PSDB; AAD35169.  
 XX  
 PT Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antibiotic-induced long QT  
 PT syndrome -  
 XX  
 PS Claim 1; Page 42; 49pp; English.  
 XX  
 CC The present invention relates to novel KCNE2 genes encoding Min-K related  
 CC (MIRP) 1 ion channel proteins and polymorphisms in these genes that are  
 CC associated with ion channel disorders including antibiotic-induced long  
 CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
 CC 57 or 116 of MIRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence

CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCNE2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human M1RP1 mutant protein (A116V).

SO Sequence 123 AA;

Query Match 99.4%; Score 628; DB 23; Length 123;  
 Best Local Similarity 99.2%; Pred. No. 5.8e-67;  
 Matches 122; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFERRIFITTYMDNMRONTTAEQALQAKVDAENFYVILYLMVIGMF 60  
 DB 1 MSTLSNFTQLEDFERRIFITTYMDNMRONTTAEQALQAKVDAENFYVILYLMVIGMF 60  
 QY 61 SFTIVAILVTVSKRREHSNDPYHOYIVEDMOEKYSQILNLESKATIHENIGAGFK 120  
 DB 61 SFTIVAILVTVSKRREHSNDPYHOYIVEDMOEKYSQILNLESKATIHENIGAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123

RESULT 13

AAB29595 standard; Protein; 123 AA.

AAB29595;

19-FEB-2001 (first entry)

Human potassium channel protein KCNE2 (M1RP1) mutant 157T.

Human: KCNE2; M1RP1; potassium channel protein; KCNE1-related;  
 Mink-related; long QT syndrome; cardiac arrhythmia;  
 drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 HERG; mutant; mutein.

Homo sapiens.  
 Synthetic.

WO200063434-A1.

26-OCT-2000.

14-APR-2000; 2000MO-US10004.

15-APR-1999; 99US-0129404.

(UTAH ) UNIV UTAH RES FOUND.

(UYVA ) UNIV YALE.

Abdott GW, Sestl F, Splawski I, Keating MT, Goldstein SAN.

WPI; 2000-672747/65.

N-PSDB; AAC64085.

Novel nucleic acids encoding M1RP1, M1RP2 and M1RP3, useful for  
 diagnosing and treating ion channel disorders, especially long QT  
 syndrome

Claim 56; Page -; 132pp; English.

The invention relates to novel ion channel proteins related to  
 KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 the invention are human and rat KCNE2 (M1RP1; AAB29585 and AAB29586,  
 respectively); human and mouse KCNE3 (M1RP2; AAB29587 and AAB29588,  
 respectively); and human and mouse KCNE4 (M1RP3; AAB29589 and AAB29590,  
 respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 potassium channels (I-KR), mutations in which are associated with long

QT syndrome. The invention also relates to methods of diagnosing long QT  
 syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 nonhuman animals comprising a heterologous ion channel protein gene  
 of the invention, a transgenic animal comprising human KCNE2 and HERG  
 DNA, and methods of and screening drugs for treating long QT syndrome  
 using KCNE2 proteins (including mutants), nucleic acids encoding them  
 and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 acids, and proteins may be used for diagnosing or treating ion channel  
 disorders, especially long QT syndrome. Transgenic animals comprising  
 KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.

The present sequence represents a mutant human KCNE2 (M1RP1)  
 specifically claimed for use in diagnostic and drug screening methods of  
 the invention.  
 Note: The present sequence is not shown in the specification, but is  
 derived from the wild-type human KCNE2 protein sequence shown on page  
 119.

Sequence 123 AA;

Query Match 99.2%; Score 627; DB 21; Length 123;  
 Best Local Similarity 99.2%; Pred. No. 7.6e-67;  
 Matches 122; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MSTLSNFTQLEDFERRIFITTYMDNMRONTTAEQALQAKVDAENFYVILYLMVIGMF 60  
 DB 1 MSTLSNFTQLEDFERRIFITTYMDNMRONTTAEQALQAKVDAENFYVILYLMVIGMF 60  
 QY 61 SFTIVAILVTVSKRREHSNDPYHOYIVEDMOEKYSQILNLESKATIHENIGAGFK 120  
 DB 61 SFTIVAILVTVSKRREHSNDPYHOYIVEDMOEKYSQILNLESKATIHENIGAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123

RESULT 14

AAB29596 standard; Protein; 123 AA.

AAB29596;

19-FEB-2001 (first entry)

Human potassium channel protein KCNE2 (M1RP1) mutant T8A.

Human: KCNE2; M1RP1; potassium channel protein; KCNE1-related;  
 Mink-related; long QT syndrome; cardiac arrhythmia;  
 drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 HERG; mutant; mutein.

Homo sapiens.  
 Synthetic.

WO200063434-A1.

26-OCT-2000.

14-APR-2000; 2000MO-US10004.

15-APR-1999; 99US-0129404.

(UTAH ) UNIV UTAH RES FOUND.

(UYVA ) UNIV YALE.

Abdott GW, Sestl F, Splawski I, Keating MT, Goldstein SAN;

WPI; 2000-672747/65.

N-PSDB; AAC64086.

Novel nucleic acids encoding M1RP1, M1RP2 and M1RP3, useful for

PT diagnosing and treating ion channel disorders, especially long QT  
 syndrome -  
 PS Claim 56; Page -, 132pp; English.  
 XX  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MiRP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (MiRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MiRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-Kr), mutations in which are associated with long  
 CC QT syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents a mutant human KCNE2 (MiRP1)  
 CC specifically claimed for use in diagnostic and drug screening methods of  
 CC the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 protein sequence shown on page  
 CC 119.  
 CC  
 SQ Sequence 123 AA;  
 Query Match 99.2%; Score 627; DB 21; Length 123;  
 Best Local Similarity 99.2%; Pred. No. 7.6e-67;  
 Matches 122; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 MSTLSNFTQLEDEYFRIFITYMDNMRONTTAEQALQAKVDANFYVILYLMVMIGMF 60  
 DB 1 MSTLSNFTQLEDEYFRIFITYMDNMRONTTAEQALQAKVDANFYVILYLMVMIGMF 60  
 QY 61 SFTIIVALTSTVSKSRREHSDPYHOYIVEDMQEKRSQILNLESKATIHENIGAAGFK 120  
 DB 61 SFTIIVALTSTVSKSRREHSDPYHOYIVEDMQEKRSQILNLESKATIHENIGAAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123  
 RESULT 15  
 AAE22097  
 ID AAE22097 standard; Protein: 123 AA.  
 XX  
 AC AAE22097;  
 XX  
 DT 25-JUL-2002 (first entry)  
 XX  
 DE Human MiRP1 mutant protein (157T).  
 XX  
 KW Human; Min-K related ion channel protein; MiRP1; ion channel disorder;  
 KW KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; muten.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT Misc-difference 57  
 XX /note- "Wild type ile substituted with Thr"  
 XX  
 PN WO200222875-A2.  
 XX  
 PD 21-MAR-2002.  
 XX

PF 11-SEP-2001; 2001WO-0528332.  
 XX  
 PR 11-SEP-2000; 2000US-231571P.  
 XX  
 PA (UYIA ) UNIV YALE.  
 XX  
 PI Goldstein SAN;  
 XX  
 DR WPI: 2002-362360/39.  
 DR N-PSDB: AAD35172.  
 XX  
 PT Novel gene encoding Min-K related ion channel protein subunit and  
 PT polymorphisms in this gene associated with antibiotic-induced long QT  
 PT syndrome -  
 XX  
 PS Claim 29; Page 46-47; 49pp; English.  
 XX  
 CC The present invention relates to novel KCNE2 genes encoding Min-K related  
 CC (MiRP) 1 ion channel proteins and polymorphisms in these genes that are  
 CC associated with ion channel disorders including antibiotic-induced long  
 CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
 CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position  
 CC encoding the amino acid positions is useful for diagnosing the presence  
 CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
 CC are useful in the development of new drug therapies which selectively  
 CC target one or more KCNE2 polymorphisms that are associated with cardiac  
 CC arrhythmias. The present sequence is human MiRP1 mutant protein (157T).  
 CC  
 SQ Sequence 123 AA;  
 Query Match 99.2%; Score 627; DB 23; Length 123;  
 Best Local Similarity 99.2%; Pred. No. 7.6e-67;  
 Matches 122; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 MSTLSNFTQLEDEYFRIFITYMDNMRONTTAEQALQAKVDANFYVILYLMVMIGMF 60  
 DB 1 MSTLSNFTQLEDEYFRIFITYMDNMRONTTAEQALQAKVDANFYVILYLMVMIGMF 60  
 QY 61 SFTIIVALTSTVSKSRREHSDPYHOYIVEDMQEKRSQILNLESKATIHENIGAAGFK 120  
 DB 61 SFTIIVALTSTVSKSRREHSDPYHOYIVEDMQEKRSQILNLESKATIHENIGAAGFK 120  
 QY 121 MSP 123  
 DB 121 MSP 123

Search completed: May 15, 2003, 14:24:43  
 Job time : 77 secs



GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:45:19 ; Search time 596.902 seconds  
(without alignments)  
10011.921 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_442

Perfect score: 369  
Sequence: 1 atgtctacttatccatctt.....ctgggtcaaatgtccccc 369

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estlin:\*  
4: em\_estnu:\*  
5: em\_estrov:\*  
6: em\_estrpl:\*  
7: em\_estro:\*  
8: em\_hlc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hlc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: gb\_gss:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_hyv:\*  
20: em\_gss\_pln:\*  
21: em\_gss\_vrt:\*  
22: em\_gss\_fun:\*  
23: em\_gss\_man:\*  
24: em\_gss\_mus:\*  
25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rtd:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	354.8	96.2	803	12	BG208163 RST27654
2	333	90.2	410	9	AI962650 q42e03.x
3	333	90.2	429	9	AI654552 AI654552
4	328.8	89.1	391	9	AI339609 q42a07.x
5	325	88.1	372	9	AI246239 q129g04.x
6	266.6	72.2	1691	11	AK008619 Mus muscu

7	257.8	69.9	470	14	D85797	D85797 Rat
8	242	65.6	1003	12	BG261965	BG261965 602373784
9	177.8	48.2	746	12	BG221966	BG221966 RST41783
10	168.2	45.6	351	13	BG938225	BG938225 AB01412
11	157	42.5	188	9	AA633404	AA633404 np69h1.s
12	119.4	32.4	121	9	AA935321	AA935321 oot1909.s
13	99	26.8	311	10	AM869303	AM869303 MR3-SM006
14	90	24.4	314	10	BB564873	BB564873 BB564873
15	70	19.0	252	14	BM783832	BM783832 K-EST0061
16	57.6	15.6	272	10	BB574249	BB574249 BB574249
17	57.2	15.5	358	10	BE486735	BE486735 174950 BA
18	53.2	14.4	716	13	BI459541	BI459541 603200548
19	52.8	14.3	464	9	AA667912	AA667912 vv19f07.x
20	52.8	14.3	603	9	AI956381	AI956381 u174e07.y
21	52.8	14.3	674	10	BB613272	BB613272 BB613272
22	52.8	14.3	754	11	AK008938	AK008938 Mus muscu
23	51.8	14.0	311	10	AM869303	AM869303 MR3-SM006
24	51	13.8	424	13	BM389584	BM389584 UI-R-CN1-
25	51	13.8	760	14	BQ194830	BQ194830 UI-R-CN1-
26	50.8	13.8	270	10	BB595946	BB595946 BB595946
27	50.8	13.8	869	12	BF540248	BF540248 602050296
28	45	12.2	986	17	CNS076KL	AL431515 T3 end of
29	42.2	11.4	484	13	BQ059218	BQ059218 BQ059218
30	42.2	11.4	687	13	BQ098508	BQ098508 BQ098508
31	42	11.4	597	13	BQ038615	BQ038615 BQ038615
32	40.4	10.9	589	13	BQ094875	BQ094875 BQ094875
33	40.4	10.9	629	13	BQ095114	BQ095114 BQ095114
34	40.4	10.9	893	14	BQ734561	BQ734561 AGENCOURT
35	39.8	10.8	309	13	BQ059046	BQ059046 BQ059046
36	38.4	10.4	634	10	AM187570	AM187570 BNGH1777
37	38.4	10.4	694	12	BF479357	BF479357 I48-3026T
38	38.4	10.4	697	13	BM301804	BM301804 MCA045A01
39	37.8	10.2	455	12	BG554741	BG554741 dac30b03.
40	37.2	10.1	539	13	BI450048	BI450048 daa97c09.
41	36.6	9.9	375	14	BM971336	BM971336 UI-CF-EC1
42	36.6	9.9	591	14	BM984243	BM984243 UI-CF-EC1
43	36.6	9.9	637	9	AL779553	AL779553 AL779553
44	36.2	9.8	431	10	BB789522	BB789522 BB789522
45	36.2	9.8	772	10	BE273157	BE273157 601142325

#### ALIGNMENTS

RESULT 1  
LOCUS BG208163 803 bp mRNA linear EST 21-APR-2001  
DEFINITION RST27654 Athersys RAGE Library Homo sapiens cDNA, mRNA sequence.  
ACCESSION BG208163  
VERSION BG208163.1 GI:13729850  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
REFERENCE  
1 (bases 1 to 803)  
Harrington, J.J., Sherf, B., Rundlett, S., Jackson, P.D., Perry, R.,  
Cain, S., Leventhal, C., Thornton, M., Ramachandran, R., Whittington, J.,  
Lerner, L., Costanzo, D., McElligott, K., Booser, S., Mays, R., Smith,  
J., Danz, J., and Ducar, M.  
Creation of genome-wide protein expression libraries using random  
activation of gene expression  
Nat. Biotechnol. 19 (5), 440-445 (2001)  
21227151  
COMMENT  
Contact: Scott J. Cain  
Athersys, Inc.  
3201 Carnegie Ave, Cleveland, OH 44115, USA  
Tel: 216 431 9900  
Fax: 216 361 9596  
Email: scai@atersys.com  
High quality sequence stop: 550.  
Location/Qualifiers



Email: cgapbs-r@mail.nih.gov  
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.  
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.  
 cDNA Library Arrayed by: Greg Lennon, Ph.D.  
 DNA Sequencing by: Washington University Genome Sequencing Center  
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: [www-bio.lnl.gov/bbrp/image/image.html](http://www-bio.lnl.gov/bbrp/image/image.html)  
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 Seq primer: -400P from Gibco  
 High quality sequence stop: 411.  
 Location/Qualifiers

## FEATURES

source

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 /clone="IMAGE:2308895"  
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 /note="Vector: p773D-Pac (Pharmacia) with a modified polylinker; Site.1: Not I; Site.2: Eco RI; Plasmid DNA from the normalized library NCI-CGAP.GC4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonoids 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo."  
 BASE COUNT 127 a 100 c 97 g 104 t 1 others  
 ORIGIN

Query Match 90.2%; Score 333; DB 9; Length 429;  
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 Matches 333; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 37 GAGCTCTCCGAAGATTTTATTTATGACAAATTTGGCGCCAGAACACAGCTG 96  
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 DB 12 GAGCTCTCCGAAGATTTTATTTATGACAAATTTGGCGCCAGAACACAGCTG 71  
 OY 97 GAGCAAGAGGCCCTCCAAAGCAAGTGTGAGAACTTCTACTATGTCATCTGTAC 156  
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 DB 72 GAGCAAGAGGCCCTCCAAAGCAAGTGTGAGAACTTCTACTATGTCATCTGTAC 131  
 OY 157 CTCATGGTATGATTTGATTTCTTCTTCATCATCTGTCATCTGTCATCTGTC 216  
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 DB 132 CTCATGGTATGATTTGATTTCTTCTTCATCATCTGTCATCTGTCATCTGTC 191  
 OY 217 AATCCAGAGAGCGGACACTTCATGACCCCTACCCACAGTATCTGTAGAGACTGG 276  
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 DB 192 AATCCAGAGAGCGGACACTTCATGACCCCTACCCACAGTATCTGTAGAGACTGG 251  
 OY 277 CAGGAAAGTACAGAGCCAAATCTTGAATCTAGAAAGTCAAGGCCACCATCATGAG 336  
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 DB 252 CAGGAAAGTACAGAGCCAAATCTTGAATCTAGAAAGTCAAGGCCACCATCATGAG 311  
 OY 337 AACATGTGTGGCGCTGGTTCAAATGTCCCCC 369  
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 DB 312 AACATGTGTGGCGCTGGTTCAAATGTCCCCC 344

RESULT 4 391 bp mRNA linear EST 29-DEC-1998  
 A1339609  
 LOCUS  
 DEFINITION qq42a07.x1 Soares\_NHMPu.S1 Homo sapiens cDNA clone IMAGE:1935156  
 3' similar to SW:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.  
 ACCESSION A1339609  
 VERSION A1339609.1 GI:4076536  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
 REFERENCE 1 (bases 1 to 391)  
 AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT  
 CONTACT: Robert Strausberg, Ph.D.  
 Email: cgapbs-r@mail.nih.gov  
 This clone is available royalty-free through LNL; contact the IMAGE Consortium ([infoimage.lnl.gov](http://infoimage.lnl.gov)) for further information.  
 Seq primer: -400P from Gibco  
 High quality sequence stop: 380.  
 Location/Qualifiers

## FEATURES

source

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 /clone="IMAGE:1935156"  
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 /tissue.type="Pooled human melanocyte, fetal heart, and pregnant uterus"  
 /lab.host="DH10B"  
 /note="Organ: mixed (see below); Vector: p773D-Pac (Pharmacia) with a modified polylinker; Site.1: Not I; Site.2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2NbHM, pregnant uterus NBHPU, and fetal heart NBH19W) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479."  
 BASE COUNT 119 a 93 c 93 g 86 t  
 ORIGIN

Query Match 89.1%; Score 328.8; DB 9; Length 391;  
 Best Local Similarity 99.4%; Pred. No. 8.8e-85;  
 Matches 330; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 38 ACGTCTCCGAAGATTTTATTTATGACAAATTTGGCGCCAGAACACAGCTG 97  
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 DB 3 ACGTCTCCGAAGATTTTATTTATGACAAATTTGGCGCCAGAACACAGCTG 62  
 OY 98 AGCAAGAGGCCCTCCAAAGCAAGTGTGAGAACTTCTACTATGTCATCTGTAC 157  
 |||||  
 DB 63 AGCAAGAGGCCCTCCAAAGCAAGTGTGAGAACTTCTACTATGTCATCTGTAC 122  
 OY 158 TCATGGTATGATTTGATTTCTTCTTCATCATCTGTCATCTGTCATCTGTC 217  
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 DB 123 TCATGGTATGATTTGATTTCTTCTTCATCATCTGTCATCTGTCATCTGTC 182  
 OY 218 AATCCAGAGAGCGGACACTTCATGACCCCTACCCACAGTATCTGTAGAGACTGG 277  
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 DB 183 AATCCAGAGAGCGGACACTTCATGACCCCTACCCACAGTATCTGTAGAGACTGG 242  
 OY 278 AGGAAAGTACAGAGCCAAATCTTGAATCTAGAAAGTCAAGGCCACCATCATGAG 337  
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 DB 243 AGGAAAGTACAGAGCCAAATCTTGAATCTAGAAAGTCAAGGCCACCATCATGAG 302  
 OY 338 ACATTTGTGTGGCGCTGGTTCAAATGTCCCCC 369  
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 DB 303 ACATTTGTGTGGCGCTGGTTCAAATGTCCCCC 334

RESULT 5 372 bp mRNA linear EST 28-JAN-1999  
 A1246239  
 LOCUS  
 DEFINITION q129g04.x1 Soares\_NHMPu.S1 Homo sapiens cDNA clone IMAGE:1857942  
 3' similar to SW:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.  
 ACCESSION A1246239  
 VERSION A1246239.1 GI:3841636

KEYWORDS	EST.
SOURCE	human.
ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
AUTHORS	NCI-CCAP <a href="http://www.ncbi.nlm.nih.gov/nciccap">http://www.ncbi.nlm.nih.gov/nciccap</a> .
TITLE	National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL	Unpublished (1997)
COMMENT	Contact: Robert Strausberg, Ph.D. Email: <a href="mailto:cgapbs-remail.nih.gov">cgapbs-remail.nih.gov</a> This clone is available royalty-free through LNL; contact the IMAGE Consortium ( <a href="mailto:info@image.llnl.gov">info@image.llnl.gov</a> ) for further information. Insert Length: 921 Std Error: 0.00 Seq primer: -40UP from GlDco High quality sequence stop: 365. Location/Qualifiers
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BASE COUNT	115 a 89 c 86 g 82 t
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Y	45 CCGAAGATTTTATTACTTATATGAGACAATTGGCGCCAGAACACACAGCTGAGCAAGA 104
DB	1 CCGAAGATTTTATTACTTATATGAGACAATTGGCGCCAGAACACACAGCTGAGCAAGA 60
Y	105 GGCCCTCCAGCCCAAAGTGTGATGCTGAGACACTTACTATGACATCGTACCTCATGGT 164
DB	61 GGCCCTCCAGCCCAAAGTGTGCTGAGAACACTTACTATGATGTCATCGTACCTCATGGT 120
Y	165 GATGATTGGAATGTTCTCTTCATCATCATGCTGGCCATCCCTGGTGAGCACTGGAATCCAA 224
DB	121 GATGATTGGAATGTTCTCTTCATCATCATGCTGGCCATCCCTGGTGAGCACTGGAATCCAA 180
Y	225 GAGACGGGACACCTCCCAATGACACCCCTACACACAGTACATTTAGAGAGACTGGCAGGAAAA 284
DB	181 GAGACGGGAAACCTCCCAATGACACCCCTACACACAGTACATTTAGAGAGACTGGCAGGAAAA 240
Y	285 GTACAAGAGCCAAATCTTGATCTAGAAAGATTCGAAGGCCACCATTCATGAGAACATTGG 344
DB	241 GTACAAGAGCCAAATCTTGATCTAGAAAGATTCGAAGGCCACCATTCATGAGAACATTGG 300
Y	345 TGGCGCTGGGTTCAAATGTCCCC 369
DB	301 TGGCGCTGGGTTCAAATGTCCCC 325
RESULT 6	
LOCUS	AK008619 1691 bp mRNA linear HTC 19-JAN-2002
DEFINITION	Mus musculus adult male stomach cDNA, RIKEN full-length enriched

ACCESSION VERSION KEYWORDS SOURCE	ORGANISM	REFERENCE AUTHORS TITLE JOURNAL MEDLINE PUBMED REFERENCE AUTHORS	TITLE JOURNAL MEDLINE PUBMED REFERENCE AUTHORS	TITLE JOURNAL MEDLINE PUBMED REFERENCE AUTHORS
AK008619.1 GI:12842913 HTC; CAP trapper. Mus musculus (strain:C57BL/6J) adult male stomach cDNA to mRNA, clone:11b;RIKEN full-length enriched mouse cDNA library clone:2200002116.	Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	Carninci, P. and Hayashizaki, Y. 1 High-efficiency full-length cDNA cloning Meth. Enzymol. 303, 19-44 (1999) 99279253 10349636 2	Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M., and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes Genome Res. 10 (10), 1617-1630 (2000) 20493374 11042159 3	Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitanai, T., Tachiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishino, T., Harada, A., Yamamoto, S., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Omata, E., Watanuki, M., Onodera, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A., and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multicapillary sequencer Genome Res. 10 (11), 1757-1771 (2000) 20530913 11076861 4
Kawaji, J., Shinagawa, A., Shibata, K., Yoshino, M., Itoh, M., Ishii, Y., Aizawa, T., Hara, A., Fukunishi, Y., Konno, H., Adachi, J., Fukuda, S., Aizawa, K., Izawa, M., Nishi, K., Kiyosawa, H., Kondo, S., Yamataka, I., Saito, T., Okazaki, Y., Gojobori, T., Bono, H., Kasukawa, T., Saito, R., Kado, K., Matsuda, H., Ashburner, M., Batalov, S., Casavant, T., Fleischmann, W., Gaasterland, T., Gissi, C., Kling, B., Kochiyama, H., Kuell, P., Lewis, S., Matsuo, Y., Nikaido, I., Pesole, G., Quackenbush, J., Schirml, L. M., Staudt, F., Suzuki, R., Tomita, M., Wagner, L., Washio, T., Sakai, K., Okido, T., Furuno, M., Aono, H., Baldarelli, R., Barish, G., Blake, J., Boffelli, D., Bojunga, N., Carninci, P., de Bonaldo, M. F., Brownstein, M. J., Bult, C., Fletcher, C., Fujita, M., Gariboldi, M., Gustincich, S., Hill, D., Hofmann, M., Hume, D. A., Kaniya, M., Lee, N. H., Lyons, P., Marchionni, L., Mashima, J., Mezzarelli, J., Mombert, P., Nordone, P., Rung, B., Ringwald, M., Rodriguez, I., Sakamoto, N., Sasaki, H., Saito, K., Schonbach, C., Seya, T., Shibata, Y., Storch, K. F., Suzuki, H., Toyko-Oka, K., Wang, K. H., Weitz, C., Whitaker, C., Wilmberg, L., Wynshaw-Boris, A., Yoshida, K., Hasegawa, Y., Kawaji, H., Kotschuki, S., and Hayashizaki, Y.				
Functional annotation of a full-length mouse cDNA collection Nature 409 (6821), 685-690 (2001) 21085660 11217851				
5 (bases 1 to 1691)				
Adachi, J., Aizawa, K., Akahira, S., Akinura, T., Aono, H., Arai, A., Arakawa, T., Baldarelli, R., Bono, H., Brownstein, M., Bult, C., Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Haneagaki, T., Hara, A., Hayatsu, N., Hill, D., Hiramoto, K., Hirooka, T., Hori, F., Hume, D., Imocant, K., Ishii, Y., Itoh, M., Izawa, M., Kasukawa, T., Kato, H., Kawaji, J., Kohima, Y., Konno, H., Konda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C., Quackenbush, J., Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, N.,				





[illegible]

RESULT	8
BG261965	
LOCUS	BG261965
DEFINITION	BG261965 1003 bp mRNA linear EST 13-FEB-2001
ACCESSION	602373778.F1 NIH_MGC_94 Mus musculus cDNA clone IMAGE:4481325 5'
VERSION	BG261965
KEYWORDS	BG261965.1 GI:12771781
SOURCE	EST.
ORGANISM	house mouse. Mus musculus

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

1 (bases 1 to 1003)  
NIH-MGC <http://mgc.ncbi.nlm.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.

Tissue Procurement: The Cepko Laboratory  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LIM)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LIM at:  
<http://image.llnl.gov>

```

FEATURES
source
Plate: L1AM10316 row: 9 column: 22
High quality sequence stop: 535.
Location/Qualifiers
1..1003
/organism="Mus musculus"

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BASE COUNT      249 a      218 c      302 g      234 t
ORIGIN
/lab_host="DH10B (phage-resistant)"
/notes=Organ: eye; Vector: PCMV-SPOF6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally; oligo-dr primed.
Average Insert size 3.3 kb. Library enriched for
full-length clones and constructed by Life Technologies.
Note: this is a NIH_MGC library."

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[illegible]

QY	241	AATACCCCTACCAACAGTACATGTGTAGAGCATGGCAGGAAAACTACAGAGCCCAATC	300
Db	331	CAGCACCCTGACCCACAGTACATCGTGGAAATTTGGCAGGAAAAAGTACAAAAAGTCAGATTC	390
QY	301	TTGAAATCTAGAAAGATAGAA--GGCCACACATCATGTAGAAACAT--TGNGCGCGCTGGGTTTC	357
Db	391	CTGCATCTGGAAAGACTCCAAAGGGCCACACATCATGTAGAAACATGGGGGGCCACCGGGTTTC	450
QY	358	AAAATGTCCCC	369
Db	451	ACAGTGTCCACC	462

RESULT 9	LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE
BG221966	BG221966	746 bp	MRNA	linear	EST 21-APR-2001	
	RT041783	Atherays	RAGE	library	Homo sapiens	CDNA, mRNA sequence.
	BG221966					
	BG221966.1	GI:13747987				
	EST.					
	human.					

REFERENCE  
AUTHORS

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.  
1 (bases 1 to 746)  
Harrington, J. J., Sherf, B., Rundlett, S., Jackson, P. D., Perry, R.,  
Cain, S., Leventhal, C., Thornton, M., Ramachandran, R., Wiltinton, J.

TITLE	Creation of genome-wide protein expression libraries using random activation of gene expression
JOURNAL	Nat. Biotechnol. 19 (5), 440-445 (2001)
MEDLINE	21227151
COMMENT	Contact: Scott J. Cain

**Athersys, Inc.**  
3201 Carnegie Ave, Cleveland, OH 44115, USA  
Tel: 216 431 9900  
Fax: 216 361 9596  
Email: [scaln@athersys.com](mailto:scaln@athersys.com)  
High quality sequence atop: 547.

```

source
1. 746
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Altersys RAGE Library"
/cell_line="Hri1080"
/notes="See 'Creation of Genome-wide Protein Expression
Libraries using Random Activation of Gene Expression',
Nature Biotechnology, In press. Note that even though the
cell type indicated is Hri1080, since a random activation
method was used, these sequence tags are not necessarily
expressed in Hri1080 under normal circumstances."
BASE COUNT      241 a      138 c      123 g      243 t      1 others
ORIGIN

```

Query Match	Best Local Match	Similarity	98.2%	Score 177.8	DB 12	Length 746
Matches 207	Conservative	0	Mismatches 13	Indels 2	Gaps 2	
Oy	1	ANGCTACTTATTCACAAATTCACACAGACGCTTGAGAGACTCTTCGAAAGATTTTATTT	60			
Ddb	526	ATGCTCACTTATTCACAAATTCACACAGACGCTTGAGAGACTCTTCGAAAGATTTTATTT	585			
Oy	61	ACTTATATGAGCAATTTGGCCGCCAGAAACACAAACGCTGAGCAGAAAGGCCCTTCACAAAGC	120			
Ddb	586	ACTTATATGAGCAATTTGGCCGCCAGAAACACAAACGCTGAGCAGAGAGGCCCTTCACAAAGC	644			
Oy	121	GTTTGATGCTGAGAACCTTCACATATGTCATCCTGTACCTCATGGTGATGATGGAATGTTTC	180			

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Db      645 GTTGATGCTGAGAACTTCTACTATGTCATCCCGAGCCCAATGANGATGTGAATGTTTC 704
      181 TCTTTCAT-CATCGTGCCATCTCTGTGAGCACTGTGAATC 221
      705 TTTTAAATAAATGGGGCATCTCTGGGAGCACTGTGAATC 746

RESULT 10
Bg38225 351 bp mRNA linear EST 11-JUN-2001
LOCUS   1AB01412 Bovine Abomasum cDNA library Bos taurus cDNA 5', mRNA
DEFINITION
ACCESSION Bg38225
VERSION    Bg38225
KEYWORDS   EST.
SOURCE     GI:14337597
ORGANISM   Bos taurus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
            Bovidae; Bovinae; Bos.
REFERENCE 1 (bases 1 to 351)
AUTHORS   Moore,S.S., Hansen,C., Li,C., Fu,A., Meng,Y. and Li,G.
TITLE      cDNA's from bovine abomasum tissue
JOURNAL    Unpublished (2001)
COMMENT    Contact: Dr. Stephen Moore
            Dept. of AENS, University of Alberta
            410 Agri/For, Dept of AENS, U of A, Edmonton, AB, T6G 2P5, Canada
            Tel: 780 492 0169
            Fax: 780 492 4265
            Email: smoores@aens.ualberta.ca
            The sequence best matches gb:AP001719 (Homo sapiens genomic DNA,
            chromosome 21q, section 63/105) in main database at high score of
            212.0 and E-value of 9e-53
PCR Primers
FORWARD: M13 Forward
BACKWARD: M13 Reverse
Seq primer: T3 primer
High quality sequence stop: 351
POLY-A-NO.

FEATURES
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            /db_xref="taxon:9913"
            /clone_11b="Bovine Abomasum cDNA library"
            /sex="Two males and one female mixed"
            /tissue_type="Gastrointestinal tissue (GIT)"
            /cell_type="Epithelial"
            /dev_stage="Young adult"
            /lab_host="XLI-BlueMRF"-strain"
            /note="Organ: Abomasum; Vector: Uni-22APXR; Site_1: EcoR
            I; Site_2: Xho I"

BASE COUNT 104 a 89 c 76 g 82 t
ORIGIN
Query Match 45.6%; Score 168.2; DB 13; Length 351;
Best Local Similarity 87.1%; Pred. No. 3.9e-38;
Matches 196; Conservative 0; Mismatches 28; Indels 1; Gaps 1;
QY 1 ATGTGACTTTTTCACAAATTCACAGAGCGTGGAGAGCGTCTCGGAGGATTTTATT 60
      127 ATGCCAATCTATTCATCTGACACAGACCTGGAAATATGCTTCAAAAGATTTTATC 186
      61 ACTAATATGACAAATTTGGCGGCGAGAACACAGACAGTGAAGAGAGCCCTCCAGCCAAA 120
      187 ACTTACATGGAAGAACTGCGACAGAGACAGCAAGCGTTGAGCAAGAGCCCTCGAAGCAAG 246
      121 GTTGATGCTGAGAACTTCTACTATGTCATCCCGAGCCCAATGANGATGTGAATGTTTC 180
      247 GTTGATGCTGAGAACTTCTACTATGTCATCCCGAGCCCAATGANGATGTGAATGTTTC 306
      181 TCTTTCAT-CATCGTGCCATCTCTGTGAGCACTGTGAATC 224

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Db      307 TCTTTCATCATCTTACGCAATCCCTGTGAGCAAGGTGAATCCAA 351
      181 TCTTTCATCATCTTACGCAATCCCTGTGAGCAAGGTGAATCCAA 351
      705 TTTTAAATAAATGGGGCATCTCTGGGAGCACTGTGAATC 746

RESULT 11
AA633404 188 bp mRNA linear EST 28-OCT-1997
LOCUS   np69h11.s1 NCI-CGAP Br2 Homo sapiens cDNA clone IMAGE:1131621.3,
DEFINITION similar to SW:MINK_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM
            CHANNEL PROTEIN 1, mRNA sequence.
ACCESSION AA633404
VERSION    AA633404
KEYWORDS   EST.
SOURCE     GI:2555264
ORGANISM   human.
            Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 188)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
            Contact: Robert Strausberg, Ph.D.
            Email: cga@bbs-rtm1.nhl.nih.gov
            Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D.
            cDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LNL at:
            www.bio.lnl.gov/bbrp/image/image.html
            Insert Length: 785 Std Error: 0.00
            Seq primer: -40m13 fwd. ET from Amersham
            High quality sequence stop: 167.

FEATURES
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            /db_xref="taxon:9606"
            /clone_11b="NCI-CGAP-Br2"
            /clone_11b="NCI-CGAP-Br2"
            /sex="female, pooled"
            /tissue_type="breast"
            /lab_host="DH10B"
            /note="Vector: pT73D-Pac (Pharmacia) with a modified
            polylinker; 1st strand cDNA was prepared from pooled bulk
            breast tumor tissue, and was then primed with a Not I -
            oligo(dT) primer. Double-stranded cDNA was ligated to Eco
            RI adaptors (Pharmacia), digested with Not I and cloned
            into the Not I and Eco RI sites of the modified pT73
            vector. This library is the normalized version of
            NCI-CGAP-Br1.1. Library was constructed by Bento Soares
            and M. Fatima Bonaldo."

BASE COUNT 66 a 45 c 45 g 32 t
ORIGIN
Query Match 42.5%; Score 157; DB 9; Length 188;
Best Local Similarity 100.0%; Pred. No. 5.4e-35;
Matches 157; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 213 TGTGAATCCCAAGAGCGGAACCTCAATGACCCCTACCAAGCTACATTTGAGAGA 272
      1 TGTGAATCCCAAGAGCGGAACCTCAATGACCCCTACCAAGCTACATTTGAGAGA 60
      273 CTGGAGAGAAAGTCAAGAGCAATCTTGAATCTAGAGATGAGGACCATCA 332
      61 CTGGAGAGAAAGTCAAGAGCAATCTTGAATCTAGAGATGAGGACCATCA 120
      333 TGAGAACATTTGGTGGCTGGGTTCAAAATGTCCCCC 365
      121 TGAGAACATTTGGTGGCTGGGTTCAAAATGTCCCCC 157

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RESULT 12  
AA935321 121 bp mRNA linear EST 07-JUL-1998  
LOCUS 007109.s1 NCI\_CGAP\_GC4 Homo sapiens cDNA clone IMAGE:1571680 3'  
DEFINITION similar to SW:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POREASSISTUM  
CHANNEL PROTEIN ; mRNA sequence.  
ACCESSION AA935321 GI:3092478  
VERSION AA935321.1  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 121)  
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgaps-remail.nih.gov  
Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael  
Emmert-Buck, M.D., Ph.D.  
CDNA Library Preparation: M. Bento Soares, Ph.D.  
CDNA Library Arrayed by: Greg Lennon, Ph.D.  
DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution Information can be  
found through the I.M.A.G.E. Consortium/Link at:  
[www.bio.livl.gov/bdrip/image/image.html](http://www.bio.livl.gov/bdrip/image/image.html)

FEATURES  
source  
1. 121  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_1ib="IMAGE:1571680"  
/clone\_1ib="NCI\_CGAP\_GC4"  
/tissue\_type="pooled germ cell tumors"  
/lab\_host="DH10B"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified  
polylinker: 1st strand cDNA was prepared from 3 pooled  
germ cell tumors, and was then primed with a Not I -  
oligo(dT) primer. Double-stranded cDNA was ligated to Eco  
RI adaptors (Pharmacia), digested with Not I and cloned  
into the Not I and Eco RI sites of the modified pT73  
vector. Library is normalized. Library was constructed by  
Bento Soares and M. Fatima Bonaldo."

BASE COUNT 33 a 31 c 28 g 29 t  
ORIGIN

Query Match 32.4%; Score 119.4; DB 9; Length 121;  
Best Local Similarity 99.2%; Pred. No. 3.8e-24;  
Matches 120; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 153 GTACCTCATGTGATGTTGGAAATGTTCTTCATCATGTCGCCATCTCGTGAGCAC 212  
|||||  
DB 1 GTACCTCATGTGATGTTGGAAATGTTCTTCATCATGTCGCCATCTCGTGAGCAC 60  
|||||

OY 213 TGTGAATCCAGAGGAGGAGACACCTGACACCCCTACACAGACAGATTGTAGAGA 272  
|||||  
DB 61 TGTGAATCCAGAGGAGGAGACACCTGACACCCCTACACAGACAGATTGTAGAGA 120  
|||||

OY 273 C 273  
DB 121 C 121

RESULT 13  
AA869303 311 bp mRNA linear EST 22-MAY-2000  
LOCUS AM869303/c  
DEFINITION MR3-SN0067-240400-006-f11 SN0067 Homo sapiens cDNA, mRNA sequence.

ACCESSION AM869303  
VERSION AM869303.1 GI:8003356  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 311)  
AUTHORS Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,  
Nagai, M.A., da Silva, M. Jr., Zago, M.A., Bordin, S., Costa, F.F.,  
Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,  
Brunstein, A., deoliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare,  
M.D., Soares, F., Brecht, R.R., Reis, L.F., de Souza, S.J. and  
Simpson, A.J.  
TITLE Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
MEDLINE 20202663  
COMMENT Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: [asimpson@ludwig.org.br](mailto:asimpson@ludwig.org.br)  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(<http://www.ludwig.org.br/scripts/gethtml2.pl?cl=6t2-MR3-SN0067-240>  
400-006-f11&t3=2000-04-24&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 11  
High quality sequence stop: 74.  
Location/Qualifiers  
1. 311  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_1ib="SN0067"  
/dev\_stage="Adult"  
/note="Organ: stomach, normal; Vector: puc18; Site: 1; Smat;  
Site: 2; Smat; A mini-library was made by cloning products  
derived from ORESTES PCR (U.S. Letters Patent application  
No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the puc 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."

BASE COUNT 60 a 80 c 76 g 95 t  
ORIGIN

Query Match 26.8%; Score 99; DB 10; Length 311;  
Best Local Similarity 90.7%; Pred. No. 4.7e-18;  
Matches 117; Conservative 0; Mismatches 10; Indels 2; Gaps 1;

OY 1 ATGTCTACTTTATTCATTTACACAGACGCTGAGAGCGTTCGAGAGATTTTAT 60  
|||||  
DB 182 ATGTCTACTTTATTCATTTACACAGACGCTGAGAGAGCTCTTCGAGAGATTTTATC 123  
|||||

OY 61 ACTTATATGACAAATGG--CGCCAGACACACACAGCCTGAGCAAGAGCCCTCAAGCCA 118  
|||||  
DB 122 ACTCATATGACAAATGGCGCGCGAGACACAGCAGTCTAGCGAGAGGCCCTCAAGCCT 63  
|||||

OY 119 AAGTGATG 127  
DB 62 AAGATGATG 54

RESULT 14  
BB564873 314 bp mRNA linear EST 29-NOV-2000  
LOCUS BB564873  
DEFINITION musculus cDNA clone 2200002116 5', mRNA sequence.  
ACCESSION BB564873 GI:11455765



Db 1 CTGATCTAGAGATCGAAGCCACCATCCATGAGACATTGTCGGGCTGGGTTCAA 60  
QY 360 AATGTCGCC 369  
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Db 61 AATGTCGCC 70

Search completed: May 21, 2003, 21:37:55  
Job time : 600.902 secs



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;; PRIOR APPLICATION NUMBER: PCT/US01/00670
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: US 60/234,687
;; PRIOR FILING DATE: 2000-09-21
;; PRIOR APPLICATION NUMBER: US 09/608,408
;; PRIOR FILING DATE: 2000-06-30
;; PRIOR APPLICATION NUMBER: US 09/774,203
;; PRIOR FILING DATE: 2001-01-29
;; NUMBER OF SEQ ID NOS: 49117
;; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
;; SEQ ID NO 33139
;; LENGTH: 372
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
;; OTHER INFORMATION: MAP TO AP000120.1
;; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.98
;; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 0.67
;; OTHER INFORMATION: EST_HUMAN HIT: A155452.1, EVALU0 0.00e+00
;; OTHER INFORMATION: SWISSPROT HIT: Q916J6, EVALU0 8.00e-67
;; OTHER INFORMATION: NT HIT: g111526220, EVALU0 0.00e+00
US-09-864-761-33139
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Query Match          100.0%; Score 369; DB 10; Length 372;
Best Local Similarity 100.0%; Pred. No. 1.3e-115;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
Db 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
QY 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
Db 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
Db 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
QY 181 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 240
Db 181 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 240
QY 241 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 300
Db 241 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 300
QY 301 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 360
Db 301 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 360
QY 361 ATGTCCCCC 369
Db 361 ATGTCCCCC 369
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RESULT 2
US-10-000-151B-5
; Sequence 5, Application US/10000151B
; Publication No. US20030013136A1
; GENERAL INFORMATION:
; APPLICANT: Balser, Jeffrey R.
; APPLICANT: George, Alfred L.
; TITLE OF INVENTION: HUMAN KRI REGULATION OF HERG POTASSIUM CHANNEL BLOCK
; FILE REFERENCE: Vanderbilt Ref No. US20030013136A1 V00120; Attorney Docket No. US2003
; CURRENT FILING DATE: 2000-10-30
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 732
; TYPE: DNA
; ORGANISM: Homo sapiens
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US-10-000-151B-5
Query Match          100.0%; Score 369; DB 9; Length 732;
Best Local Similarity 100.0%; Pred. No. 2e-115;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
Db 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
QY 74 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 133
Db 74 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 133
QY 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
Db 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
QY 134 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 193
Db 134 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 193
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
Db 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
QY 194 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 253
Db 194 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 253
QY 181 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 240
Db 181 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 240
QY 254 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 313
Db 254 TCTTTTCATCATCGTGGCCATCTCTGTGAGACCTGTGAATCCAAAGAGGGAACACTCC 313
QY 241 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 300
Db 241 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 300
QY 314 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 373
Db 314 AATGACCCCTACACAGCTACATGTAGAGAGCTGGGAGGAGAAAGTCAAGAGCCAAATC 373
QY 301 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 360
Db 301 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 360
QY 374 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 433
Db 374 TTGAATCTAGAGAAGATGAGAGGCGCCATCCATGAGAAATTTGGTGGGCTGTTCAA 433
QY 361 ATGTCCCCC 369
Db 434 ATGTCCCCC 442
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RESULT 3
US-10-227-195A-1
; Sequence 1, Application US/10227195A
; Publication No. US20030077633A1
; GENERAL INFORMATION:
; APPLICANT: Cox, David
; APPLICANT: Arnold, Deana
; TITLE OF INVENTION: Haplotype structure of chromosome 21
; FILE REFERENCE: 103001
; CURRENT APPLICATION NUMBER: US/10/227,195A
; CURRENT FILING DATE: 2002-11-18
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 113604
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7175..7204, 36973, 66372, 76921, 81512, 88727
; OTHER INFORMATION: n = G or C
US-10-227-195A-1
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Query Match          100.0%; Score 369; DB 9; Length 113604;
Best Local Similarity 100.0%; Pred. No. 3.6e-114;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
Db 1 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 60
QY 17476 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 17535
Db 17476 ATGCTACTTTATTCATTTTCACACAGACGCTGGAAGAGCTCTTCCGAAGATTTTATT 17535
QY 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
Db 61 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 120
QY 17536 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 17595
Db 17536 ACTTATATGAGCAATTTGGCGCCAGACACAGACAGCTGTGGAAGAGGCGCTCCAGCCAAA 17595
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
Db 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 180
QY 17596 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 17655
Db 17596 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATGATGATGATGTC 17655
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OY 181 TCTTTCATCGTGGCCATCTGTGAGCAGTGTGAATCCAAAGAGCGGGAACACTCC 240
      |||||||
Db 17656 TCTTTCATCGTGGCCATCTGTGAGCAGTGTGAATCCAAAGAGCGGGAACACTCC 17715

OY 241 AATGACCCCTACACACAGTACATTTGAGAGCTGGCAGGAAAAAGTACAAGACCAATC 300
      |||||||
Db 17716 AATGACCCCTACACACAGTACATTTGAGAGCTGGCAGGAAAAAGTACAAGACCAATC 17775

OY 301 TTGAATCTAGAGAAGATGGAAGGCCACCATCATGAGAACATTTGGTGGGCTGGGTTCAAA 360
      |||||||
Db 17776 TTGAATCTAGAGAAGATGGAAGGCCACCATCATGAGAACATTTGGTGGGCTGGGTTCAAA 17835

OY 361 ATGTCCCCC 369
      |||||||
Db 17836 ATGTCCCCC 17844

RESULT 4
US-10-227-195A-2
; Sequence 2, Application US/10227195A
; Publication No. US20030077633A1
; GENERAL INFORMATION:
; APPLICANT: Cox, David
; APPLICANT: Arnold, Deana
; TITLE OF INVENTION: Haplotype structure of chromosome 21
; FILE REFERENCE: 103001
; CURRENT APPLICATION NUMBER: US/10/227,195A
; NUMBER OF SEQ ID NOS: 2
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 113604
; TYPE: DNA
; ORGANISM: Human
US-10-227-195A-2

Query Match 100.0%; Score 369; DB 9; Length 113604;
Best Local Similarity 100.0%; Pred. No. 3.6e-114;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 AATGCTACTTTATTCATTTTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 60
      |||||||
Db 17476 AATGCTACTTTATTCATTTTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 17535

OY 61 ACTTATTTGACATTTGGCCCGCAGAACACACAGCTGAGCAAGAGCCCTCCAAAGCAA 120
      |||||||
Db 17536 ACTTATTTGACATTTGGCCCGCAGAACACACAGCTGAGCAAGAGCCCTCCAAAGCAA 17595

OY 121 GTTGATGCTGAGAACTTCTACTATGTGATCTGTACCTCATGTGATGATTTGAT 180
      |||||||
Db 17596 GTTGATGCTGAGAACTTCTACTATGTGATCTGTACCTCATGTGATGATTTGAT 17655

OY 181 TCTTTCATCGTGGCCATCTGTGAGCAGTGTGAATCCAAAGAGCGGGAACACTCC 240
      |||||||
Db 17656 TCTTTCATCGTGGCCATCTGTGAGCAGTGTGAATCCAAAGAGCGGGAACACTCC 17715

OY 241 AATGACCCCTACACACAGTACATTTGAGAGCTGGCAGGAAAAAGTACAAGACCAATC 300
      |||||||
Db 17716 AATGACCCCTACACACAGTACATTTGAGAGCTGGCAGGAAAAAGTACAAGACCAATC 17775

OY 301 TTGAATCTAGAGAAGATGGAAGGCCACCATCATGAGAACATTTGGTGGGCTGGGTTCAAA 360
      |||||||
Db 17776 TTGAATCTAGAGAAGATGGAAGGCCACCATCATGAGAACATTTGGTGGGCTGGGTTCAAA 17835

OY 361 ATGTCCCCC 369
      |||||||
Db 17836 ATGTCCCCC 17844

RESULT 5
US-09-864-761-20233
; Sequence 20233, Application US/09864761
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; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aecmlca-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 20233
; LENGTH: 312
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000052.1
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HB1100, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.92
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.2
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
; OTHER INFORMATION: EST_HUMAN HIT: A1246239.1, EVALU0 0.00e+00
; OTHER INFORMATION: SWISSPROT HIT: Q9Y6J6, EVALU0 3.00e-55
; OTHER INFORMATION: NT HIT: AF302095.1, EVALU0 0.00e+00
US-09-864-761-20233

Query Match 84.6%; Score 312; DB 10; Length 312;
Best Local Similarity 100.0%; Pred. No. 3.4e-96;
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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OY 54 TTTTATTACTTATATGGAATTTGGCCGACAGACAGCTGAGAGAGGCGCTTCCA 113
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Db 1 TTTTATTACTTATATGGAATTTGGCCGACAGACAGCTGAGAGAGGCGCTTCCA 60
OY 114 AGCGAAGTTGATGCTGAGAACTCTCTACTATGTCATCTGCTGACTGCTGAGTATGG 173
      |||||||
Db 61 AGCGAAGTTGATGCTGAGAACTCTCTACTATGTCATCTGCTGACTGCTGAGTATGG 120
OY 174 AATGTCCTTTCATCATCTGCGCCATCTGCTGAGCAGCTGTGAATTCAGAGCGGA 233
      |||||||
Db 121 AATGTCCTTTCATCATCTGCGCCATCTGCTGAGCAGCTGTGAATTCAGAGCGGA 180
OY 234 ACACTCCAAATGACCCCTTACACCACTGTCATTTGTAGAGACTGGCAGGAAAGTACAGAG 293
      |||||||
Db 181 ACACTCCAAATGACCCCTTACACCACTGTCATTTGTAGAGACTGGCAGGAAAGTACAGAG 240
OY 294 CCAATCTTGAAATCTAGAGAAATGGAAGGCGCACCATCATGAGAACTGGTGGGCTGG 353
      |||||||
Db 241 CCAATCTTGAAATCTAGAGAAATGGAAGGCGCACCATCATGAGAACTGGTGGGCTGG 300
OY 354 GTTCAAAATGTC 365
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Db 301 GTTCAAAATGTC 312
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RESULT 6
US-09-864-761-3463
; Sequence 3463, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aeonica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263,6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
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; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 3463
; LENGTH: 450
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000052.1
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.3
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 0.92
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.2
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.94
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 0.88
US-09-864-761-3463
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Query Match 79.9% Score 295; DB 10; Length 450;
Best Local Similarity 100.0%; Pred. No. 2.6e-90;
Matches 295; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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OY 1 ATGCTACTTTATCATATTTACACAGACGCTGGAAGAGCTCTCCGAAGATTTTATT 60
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Db 156 ATGCTACTTTATCATATTTACACAGACGCTGGAAGAGCTCTCCGAAGATTTTATT 215
OY 61 ACTTATATGACATTTGGCGCCGAGAACACACAGCTGTGAGCAGAGAGGCGCTCCAGCCAA 120
      |||||||
Db 216 ACTTATATGACATTTGGCGCCGAGAACACACAGCTGTGAGCAGAGAGGCGCTCCAGCCAA 275
OY 121 GTGAGCGAGAGACTGTACTATGTCACTGCTGACTCATGTGATGATGGAATGTC 180
      |||||||
Db 276 GTGAGCGAGAGACTGTACTATGTCACTGCTGACTCATGTGATGATGGAATGTC 335
OY 181 TCTTTCATCATCTGCGCCATCTGCTGAGCAGCTGTGAATTCAGAGAGGAGACACTCC 240
      |||||||
Db 336 TCTTTCATCATCTGCGCCATCTGCTGAGCAGCTGTGAATTCAGAGAGGAGACACTCC 395
OY 241 AATGACCCCTTACACCACTGATTTGTAGAGAGCTGGCAGGAAAGTACAGAGCC 295
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Db 396 AATGACCCCTTACACCACTGATTTGTAGAGAGCTGGCAGGAAAGTACAGAGCC 450
RESULT 7
US-09-864-761-16671
; Sequence 16671, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO
; FILE REFERENCE: Aeonica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263,6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
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; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 16671
; LENGTH: 471
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000120.1
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.98
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 0.67
; US-09-864-761-16671

Query Match          63.1%; Score 233; DB 10; Length 471;
Best Local Similarity 100.0%; Pred. No. 3.8e-69;
Matches 233; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 239 AGTCTACTTATTCACAAATTTACACAGACGCTGGAAGAGCTTCCGAAGATTTTAT 298
QY 61 ACTTATATGACAAATTTGGCCGACAAACACACAGCTGACCAAGAGCCCTCCAGCCAAA 120
DB 299 ACTTATATGACAAATTTGGCCGACAAACACACAGCTGACCAAGAGCCCTCCAGCCAAA 358
QY 121 GTTGATCGGAGAACTTCTACTATGTATGTCCTGATCCCTGATGATGATGATGATGAT 180
DB 359 GTTGATCGGAGAACTTCTACTATGTATGTCCTGATCCCTGATGATGATGATGATGATG 418
QY 181 TCTTTCATCATCGTGGCCATCTGCTGAGACACTGTGAATCCAAAGAGAGGGA 233
DB 419 TCTTTCATCATCGTGGCCATCTGCTGAGACACTGTGAATCCAAAGAGAGGGA 471

RESULT 8
US-09-864-761-20783/C
; Sequence 20783, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aemolca-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
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; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 20783
; LENGTH: 231
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AP000121.1
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.1
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 0.94
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 0.59
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 0.74
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.66
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 0.66
; OTHER INFORMATION: SWISSPROT HIT: P15382, EVALUATE 2.00e-39
; OTHER INFORMATION: EST_HUMAN HIT: AW847275.1, EVALUATE 5.60e-01
; OTHER INFORMATION: NT HIT: AF135188.1, EVALUATE 1.00e-127
; US-09-864-761-20783

Query Match          14.4%; Score 53.2; DB 10; Length 231;
Best Local Similarity 63.6%; Pred. No. 5.6e-08;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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DB 207 CCTTACGCTCTCATGATGATGATGATGATGATGATGATGATGATGATGATGATG 148
QY 210 CACTGTGAATCCAGAGAGAGGGAACCTCCAAATGACCCCTACACCAAGTATG--T 266
DB 147 CTACATCGGCTCCAGAGAGGAGCTGAGACACTCGAAGAGCCCATTTCAACGCTACATGAGTC 88
QY 267 AGAGAGCTGGCAGGAAAGTACAGAGAGCAATC 300
DB 87 CGATCGCTGGCAAGAGAGAGGAGCAAGGCTATGTC 54

RESULT 9
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GenCore version 5.1.4.p5.4578  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:16:44 ; Search time 89,8202 Seconds

(without alignments)  
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Scoring table: IDENTITY\_NDC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
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3	369	100.0	372	22	AA500245	Human potassium ch
4	369	100.0	471	22	AAAF80269	Nucleotide sequenc
5	369	100.0	600	22	ABA09192	Human MIRP1 homolo
6	369	100.0	600	22	AAK52645	Human polynucleoti
7	369	100.0	655	22	AAK51661	Human polynucleoti
8	369	100.0	732	21	AAAC64071	Human potassium ch
9	369	100.0	732	24	ABK86573	cDNA encoding huma

10	369	100.0	732	24	AAAD35170	Human KCNE2 wild t
11	367.4	99.6	732	21	AAAC64083	Human potassium ch
12	367.4	99.6	732	21	AAAC64084	Human potassium ch
13	367.4	99.6	732	21	AAAC64085	Human potassium ch
14	367.4	99.6	732	21	AAAC64086	Human potassium ch
15	367.4	99.6	732	24	AAAD35169	Human KCNE2 mutant
16	367.4	99.6	732	24	AAAD35171	Human KCNE2 mutant
17	367.4	99.6	732	24	AAAD35172	Human KCNE2 mutant
18	367.4	99.6	732	24	AAAD35173	Human KCNE2 mutant
19	312	84.6	312	22	ABAA4938	Human breast cell
20	312	84.6	312	22	ABAA4939	Human foetal liver
21	312	84.6	312	22	ABAA4940	Human foetal liver
22	312	84.6	312	22	ABAA4941	Human foetal liver
23	312	84.6	312	22	ABAA4942	Human foetal liver
24	312	84.6	312	22	ABAA4943	Human foetal liver
25	312	84.6	312	22	ABAA4944	Human foetal liver
26	312	84.6	312	22	ABAA4945	Human foetal liver
27	312	84.6	312	22	ABAA4946	Human foetal liver
28	295	79.9	450	22	ABAA4947	Human foetal liver
29	295	79.9	450	22	ABAA4948	Human foetal liver
30	295	79.9	450	22	ABAA4949	Human foetal liver
31	295	79.9	450	22	ABAA4950	Human foetal liver
32	295	79.9	450	22	ABAA4951	Human foetal liver
33	295	79.9	450	22	ABAA4952	Human foetal liver
34	295	79.9	450	22	ABAA4953	Human foetal liver
35	295	79.9	450	22	ABAA4954	Human foetal liver
36	295	79.9	450	22	ABAA4955	Human foetal liver
37	263.4	71.4	372	22	ABAA4956	Human foetal liver
38	263.4	71.4	372	22	ABAA4957	Human foetal liver
39	233	63.1	471	22	ABAA4958	Human foetal liver
40	233	63.1	471	22	ABAA4959	Human foetal liver
41	53.8	14.6	65	24	ABAA4960	Human foetal liver
42	53.2	14.4	231	22	ABAA4961	Human foetal liver
43	53.2	14.4	231	22	ABAA4962	Human foetal liver
44	53.2	14.4	231	22	ABAA4963	Human foetal liver
45	53.2	14.4	231	22	ABAA4964	Human foetal liver

## ALIGNMENTS

RESULT 1	AA124432	standard; DNA; 372 BP.
ID	AA124432	standard; DNA; 372 BP.
AC	AA124432	
XX		
DT	12-OCT-2001	(first entry)
DE	Probe #14365 for gene expression analysis in human cervical cell sample.	
XX		
KW	Probe; human; microarray; gene expression; cervical epithelial cell;	
KW	cervical cancer; ss.	
OS	Homo sapiens.	
XX		
PN	W0200157278-A2.	
XX		
PD	09-AUG-2001.	
XX		
PF	30-JAN-2001; 2001WO-US00670.	
XX		
PR	04-FEB-2000; 2000US-0180312.	
XX		
PR	26-MAY-2000; 2000US-0207456.	
XX		
PR	30-JUN-2000; 2000US-0808408.	
XX		
PR	03-AUG-2000; 2000US-0632366.	
XX		
PR	21-SEP-2000; 2000US-0234687.	
XX		
PR	27-SEP-2000; 2000US-0236359.	
XX		
PR	04-OCT-2000; 2000GB-0024263.	
XX		
PA	(MOLE-) MOLECULAR DYNAMICS INC.	
XX		
PI	Penn SG, Hanzel DK, Chen W, Rank DR;	

XX WIPI: 2001-488901/53.  
DR Human genome-derived single exon nucleic acid probes useful for  
XX analyzing gene expression in human cervical epithelial cells -  
PT  
XX  
PS Claim 25; SEQ ID No 14365; 487pp; English.  
XX  
CC The present invention relates to human single exon nucleic acid probes  
CC (SENPs). The present sequence is one such probe. The SENPs are derived  
CC from human Hela cells. The SENPs can be used to produce a single exon  
CC microarray, which can be used for measuring human gene expression in a  
CC sample derived from human cervical epithelial cells. By measuring gene  
CC expression, the probes are therefore useful in grading and/or staging  
CC of diseases of the cervix, notably cervical cancer.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;  
Query Match 100.0%; Score 369; DB 22; Length 372;  
Best Local Similarity 100.0%; Pred. No. 2,4e-101;  
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 ATGCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAGAGATTTTATT 60  
DB 1 ATGCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAGAGATTTTATT 60  
QY 61 ACTTATATGACAAATTTGGCCGACACACAAACAGCTGAGAGAGGCTCCCAAGCCAAA 120  
DB 61 ACTTATATGACAAATTTGGCCGACACACAAACAGCTGAGAGAGGCTCCCAAGCCAAA 120  
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATTTGAATGTC 180  
DB 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATTTGAATGTC 180  
QY 181 TCTTTCATCATGCTGGCCATCTCTGTGAGCACTGTGAATCCAAAGAGGGAACCTCC 240  
DB 181 TCTTTCATCATGCTGGCCATCTCTGTGAGCACTGTGAATCCAAAGAGGGAACCTCC 240  
QY 241 AATGACCCCTACACAGTACATGTGAGAGCTGGAGGAAAGTCAAGAGCCAAATC 300  
DB 241 AATGACCCCTACACAGTACATGTGAGAGCTGGAGGAAAGTCAAGAGCCAAATC 300  
QY 301 TTGAATCTAGAAAGATGGAAGGCGACCATCATGAGAAATTTGGTGGGCTGGTCAAA 360  
DB 301 TTGAATCTAGAAAGATGGAAGGCGACCATCATGAGAAATTTGGTGGGCTGGTCAAA 360  
QY 361 ATGTCCCCC 369  
DB 361 ATGTCCCCC 369  
RESULT 2  
AAI09965  
ID AAI09965 standard; DNA: 372 BP.  
XX  
XX AAI09965;  
XX  
XX 09-OCT-2001 (first entry)  
XX  
DE Probe #9956 used to measure gene expression in human breast sample.  
XX  
XX Probe: human; breast disease; breast cancer; development disorder; ss;  
XX Inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
XX  
XX Homo sapiens.  
XX  
XX MO200157270-A2.  
XX  
XX 09-AUG-2001.  
XX

PF 29-JAN-2001; 2001MO-US00661.  
XX  
XX 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
XX WIPI: 2001-476286/51.  
XX  
XX Novel single exon nucleic acid probe used to measuring gene expression  
XX in a human breast -  
XX  
XX Claim 25; SEQ ID No 9956; 322pp; English.  
XX  
XX The present invention relates to novel single exon nucleic acid probes.  
XX The present sequence is one such probe. The probes are useful for  
XX measuring human gene expression in a human breast sample, where the probe  
XX hybridises at high stringency to a nucleic acid expressed in the human  
XX breast. The probes are useful for predicting, diagnosing, grading,  
XX staging, monitoring and prognosing diseases of the human breast,  
XX particularly those diseases with polygenic aetiology. The diseases  
XX include: breast cancer, disorders of development, inflammatory diseases  
XX of the breast, fibrocystic changes, proliferative breast disease and  
XX non-carcinoma tumours.  
XX Note: The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;  
Query Match 100.0%; Score 369; DB 22; Length 372;  
Best Local Similarity 100.0%; Pred. No. 2,4e-101;  
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 ATGCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAGAGATTTTATT 60  
DB 1 ATGCTACTTTATTCACACAGACGCTGGAAGAGCTTCCGAGAGATTTTATT 60  
QY 61 ACTTATATGACAAATTTGGCCGACACACAAACAGCTGAGAGAGGCTCCCAAGCCAAA 120  
DB 61 ACTTATATGACAAATTTGGCCGACACACAAACAGCTGAGAGAGGCTCCCAAGCCAAA 120  
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATTTGAATGTC 180  
DB 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGATGATTTGAATGTC 180  
QY 181 TCTTTCATCATGCTGGCCATCTCTGTGAGCACTGTGAATCCAAAGAGGGAACCTCC 240  
DB 181 TCTTTCATCATGCTGGCCATCTCTGTGAGCACTGTGAATCCAAAGAGGGAACCTCC 240  
QY 241 AATGACCCCTACACAGTACATGTGAGAGCTGGAGGAAAGTCAAGAGCCAAATC 300  
DB 241 AATGACCCCTACACAGTACATGTGAGAGCTGGAGGAAAGTCAAGAGCCAAATC 300  
QY 301 TTGAATCTAGAAAGATGGAAGGCGACCATCATGAGAAATTTGGTGGGCTGGTCAAA 360  
DB 301 TTGAATCTAGAAAGATGGAAGGCGACCATCATGAGAAATTTGGTGGGCTGGTCAAA 360  
QY 361 ATGTCCCCC 369  
DB 361 ATGTCCCCC 369  
RESULT 3  
AAS00245  
ID AAS00245 standard; DNA: 372 BP.  
XX



```
XX AC AAS00245;
XX XX
XX DT 10-MAY-2001 (first entry)
XX DE Human potassium channel regulatory protein, Mink2, DNA sequence.
XX XX
XX KM Human: Mink2; potassium channel; cardiac arrhythmia; hypertension; ds;
XX KM angina; asthma; diabetes; renal insufficiency; urinary incontinence;
XX KM irritable colon; epilepsy; cerebrovascular ischaemia; autoimmune disease.
XX OS Homo sapiens.
XX XX
XX FH Key Location/Qualifiers
XX FT CDS 1..372
XX FT /tag= a
XX FT /product= "MINK2 potassium channel protein"
XX XX
XX PN MO200114403-A1.
XX XX
XX PD 01-MAR-2001.
XX XX
XX PF 18-AUG-2000; 2000MO-US22799.
XX XX
XX PR 20-AUG-1999; 99US-0379201.
XX XX
XX PA (UYCA-) UNIV CASE WESTERN RESERVE.
XX XX
XX PI Flicker E, Wible B, Brown AM;
XX XX
XX DR WPI: 2001-218424/22.
XX DR P-PSDB; AAB00215.
XX XX
XX PT Novel potassium channel gene termed Mink2 encoding potassium channel
XX PT regulatory protein, useful for screening compounds that are useful for
XX PT treating diseases caused by aberrant potassium activity -
XX XX
XX PS Claim 1; Fig 9; 39pp; English.
XX XX
XX CC The sequence represents the coding sequence of human potassium channel
XX CC regulatory protein, Mink2. Mink2 sequence is useful for producing a
XX CC potassium channel regulatory protein useful for in vitro or in vivo
XX CC screening of agonistic or antagonistic compounds that are useful for
XX CC treating diseases caused by aberrant potassium activity, such as human
XX CC cardiac arrhythmias, hypertension, angina, asthma, diabetes, renal
XX CC insufficiency, urinary incontinence, irritable colon, epilepsy,
XX CC cerebrovascular ischaemia, and autoimmune disease.
XX XX
XX SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;
XX XX
XX Query Match 100.0%; Score 369; DB 22; Length 372;
XX Best Local Similarity 100.0%; Pred. No. 2.4e-101;
XX Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX XX
XX QY 1 ATGCTACTTATTCACATTTTCACACAGACGCTGGAAAGCGTCTCCGAGGATTTTAT 60
XX DB 1 ATGCTACTTATTCACATTTTCACACAGACGCTGGAAAGCGTCTTCGAGGATTTTAT 60
XX QY 61 ACTATATGACAAATTTGGCGCCAGACAAACAGCTGAGCAAGAGCGCTCCAAAGCCAAA 120
XX DB 61 ACTATATGACAAATTTGGCGCCAGACAAACAGCTGAGCAAGAGCGCTCCAAAGCCAAA 120
XX QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACTCTCATGTGATGATTTGATGTC 180
XX DB 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACTCTCATGTGATGATTTGATGTC 180
XX QY 181 TCTTTCATTCATCGTGGCAGTCCTGTGAGACGTGAAATCCAAAGAGCGGAACACTCC 240
XX DB 181 TCTTTCATTCATCGTGGCAGTCCTGTGAGACGTGAAATCCAAAGAGCGGAACACTCC 240
XX QY 241 AATGACCCCTACACAGTACATTTGTAGAGAGACTGGCAGGAAAAGTCAAGAGCCAAATC 300
XX DB 241 AATGACCCCTACACAGTACATTTGTAGAGAGACTGGCAGGAAAAGTCAAGAGCCAAATC 300
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```
QY 301 TTGAATCTAGAGAAATCGAAGGCCACCATTCATGAGAACATTGNGCGGTGTTCAA 360
DB 301 TTGAATCTAGAGAAATCGAAGGCCACCATTCATGAGAACATTGNGCGGTGTTCAA 360
QY 361 ATGTCCCCC 369
DB 361 ATGTCCCCC 369
XX XX
XX RESULT 4
XX ID AAF80269
XX ID AAF80269 standard; DNA; 471 BP.
XX XX
XX AC AAF80269;
XX XX
XX DT 29-JUN-2001 (first entry)
XX DE Nucleotide sequence of human potassium channel subunit Isk2.
XX XX
XX KM Human; potassium channel; Isk2; gene therapy; gastric motility;
XX KM gastric acid secretion; anti-arrhythmic agent; myocardial infarction; ss.
XX OS Homo sapiens.
XX XX
XX FH Key Location/Qualifiers
XX FT CDS 79..450
XX FT /tag= a
XX FT /product= "potassium channel subunit Isk2"
XX XX
XX PN MO200127246-A1.
XX XX
XX PD 19-APR-2001.
XX XX
XX PF 10-OCT-2000; 2000MO-US28014.
XX XX
XX PR 12-OCT-1999; 99US-0158781.
XX XX
XX PA (MERT ) MERCK & CO INC.
XX XX
XX PI Swanson RJ, Liu Y, Folander K;
XX XX
XX DR WPI: 2001-273764/28.
XX DR P-PSDB; AAB67800.
XX XX
XX PT New DNA encoding the Isk2 potassium channel subunit, useful e.g. for
XX PT detecting mutations and screening for therapeutic agents -
XX XX
XX PS Claim 3; Fig 1A; 46pp; English.
XX XX
XX CC The present sequence encodes a human potassium channel subunit,
XX CC designated Isk2. The Isk2 polynucleotide, and derived probes, are
XX CC used diagnostically to detect mutations in the Isk2 gene, to determine
XX CC levels of mRNA expression and to isolate homologous sequences; for
XX CC recombinant expression of Isk2; in gene therapy to increase potassium
XX CC channel activity and to generate transgenic animals, as models and
XX CC for drug screening. Recombinant Isk2 is used for studying biochemical
XX CC activity of Isk2 and its role in disorders of gastric motility and
XX CC gastric acid secretion, and to raise specific antibodies. Isk2
XX CC modulators are potentially useful for treating diseases associated with
XX CC increased or reduced potassium channel activity, e.g. as
XX CC anti-arrhythmic agents for treating myocardial infarction and as
XX CC regulators of gastric acid secretion.
XX XX
XX SQ Sequence 471 BP; 143 A; 110 C; 103 G; 115 T; 0 other;
XX XX
XX Query Match 100.0%; Score 369; DB 22; Length 471;
XX Best Local Similarity 100.0%; Pred. No. 2.7e-101;
XX Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX XX
XX QY 1 ATGTCTACTTATTCACATTTTCACACAGACGCTGGAAGAGCTCTCCGAGGATTTTAT 60
XX DB 79 ATGTCTACTTATTCACATTTTCACACAGACGCTGGAAGAGCTCTCTCGAAGATTTTAT 138
```

Oy	61	ACTTATATGACCAATTGGCGCCACACACACAAACACTGTAGCAAGAGGCCCTCCAAAGCCAAA	120
Db	139	ACTTATATGACCAATTGGCGCCACACACACAAACACTGTAGCAAGAGGCCCTCCAAAGCCAAA	138
Oy	121	GTTGATGCTGGAACCTTCTACTATGTCATCCTGTACCTCATGCTGTGATGATTTGAATGTTTC	180
Db	199	GTTGATGCTGGAACCTTCTACTATGTCATCCTGTACCTCATGCTGTGATGATTTGAATGTTTC	258
Oy	181	TCTTTCATCATCATGAGGCCCATCTGTGTGACACTGTGAAATCCAAAGATAGGGGAACACTCC	240
Db	259	TCTTTCATCATCATGAGGCCCATCTGTGTGACACTGTGAAATCCAAAGATAGGGGAACACTCC	318
Oy	241	AATGACCCCTACCAACAGTACATTGTAGAGACTGTGCAGGAAAAAGTACCAAGAGCCCAATC	300
Db	319	AATGACCCCTACCAACAGTACATTGTAGAGACTGTGCAGGAAAAAGTACCAAGAGCCCAATC	378
Oy	301	TTGATCTAGAAAGATGGAAGCGCACCATCATATAGAAACATTGGTGCGGCTGGGTTCCAA	360
Db	379	TTGATCTAGAAAGATGGAAGCGCACCATCATATAGAAACATTGGTGCGGCTGGGTTCCAA	438
Oy	361	ATGTCCCC 369	
Db	439	ATGTCCCC 447	

RESULT 5  
ABA09192  
ID ABA09192 standard; cDNA; 600 BP.

DT 11-JAN-2002 (first entry)

DE Human MIRP1 homologue-encoding cDNA, SEQ ID NO:968.

KM Human:tokine: cell proliferation: cell differentiation: growth factor:  
KM hematopoiesis regulation: tissue growth: immunomodulator: actvlin:  
KM inhibin: chemokaxis: chemokinesis: chromolysis: oncogenesis:  
KM proliferation: metastasis: cancer: tumour: haematopoietic disorder:  
KM myeloid cell disorder: lymphoid cell disorder: asthma: arthritis:  
KM chronic inflammatory condition: proliferative retinopathy:  
KM atherosclerosis: coronary heart disease: arterial ischaemia:  
KM bone disorder: osteoporosis: vascular growth disorder:  
KM tissue regeneration: wound healing: infection: immune disorder:  
KM cell culture: drug screening: gene therapy: antiinflammatory:  
KM antiaesthetic: antiarthritis: haemostatic: antiarteriosclerotic:  
KM cytostatic: osteoponthic: vasotropic: cardiant: virucide: antibacterial:  
KM antifungal: vulnary: antilacer: ss.

**OS Homo sapiens.**

PN W0200157188-A2.

PD 09-AUG-2001.

PF 05-FEB-2001; 2001WO-US03800.

PR 03-FEB-2000; 2000US-0496914.

[illegible]

PA (HYSE-) HYSEQ INC.

Tang YT, Liu C, Drmanac RT:

WPI: 2001-457740/49

DK P-PSUB; ABB11948  
XX

PT Human proteins and DNA encoding sequences useful for preventing,  
PT treating or ameliorating a medical condition in a mammalian subject  
PT e.g. arthritis and cancer -

xx Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and sequences ABA08225-ABA09574 represent nucleic acids encoding them. The invention also relates to vectors and recombinant host cells comprising a nucleotide of the invention, methods of producing the novel polypeptides, antibodies against the polypeptides, methods of detecting the nucleosides or polypeptides in a sample, and methods of identifying compounds which bind to polypeptides of the invention. Although novel, many of the polypeptides of the invention have homology to known proteins, thereby giving an insight into their probable biological activities, and hence potential therapeutic applications. The polypeptides of the invention may have various activities, including cytokine, cell proliferation or cell differentiation activities; stem cell growth factor activity; haematopoietic regulatory activity; tissue growth activity; immunomodulatory activity; activin- or inhibin-related activities; chemotactic or chemokinetic activities; haemostatic, thrombotic or thrombolytic activities; receptor or ligand activities; or may be involved in oncogenesis, cancer cell proliferation or metastasis. Depending on their biological activities, polypeptides and nucleosides of the invention are useful for preventing, treating or ameliorating medical conditions, e.g., by protein or gene therapy. Such conditions include cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell disorders), chronic inflammatory conditions (e.g., asthma or arthritis), proliferative retinopathy, atherosclerosis, coronary heart disease, arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal vascular growth. Polypeptides involved with tissue regeneration and repair (or nucleic acids encoding them) may be used to promote wound healing (e.g., of burns, incisions and ulcers), while those with immunomodulatory activities may be used in the treatment of viral, bacterial and fungal infections in addition to immune disorders. Polypeptides with growth factor activity may be used in cell cultures to promote cell growth. For example, such polypeptides may be used to manipulate stem cells in culture to give rise to neuroepithelial cells that can be used to augment or replace cells damaged by illness, autoimmune disease or accidental damage. The polypeptides and nucleosides may also be used in the diagnosis of the above conditions, and in drug screening techniques. The present sequence represents a cDNA encoding a novel human polypeptide of the invention.

**SQ** Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;

Query Match	100.0%	Pred. 369; DB 22,	Length 600;
Best Local Similarity	100.0%;	Score. No. 3e-101;	
Matches 369; Conservative	0;	Mismatches 0;	Indels 0; Gaps 0.

OY	1	ATGTTACTTTCATTCATTTTCACAGAGCGTGGAAAGAGTCTTCGGAAGATTTTAT	60
Db	38	ATGTTACTTTCATTCATTTTCACAGAGCGTGGAAAGAGTCTTCGGAAGATTTTAT	97
OY	61	ACTTATATGAGCAATTTGGCGCCAGAACACACAGCTGAGCAAGAGGCCCTCCAAAGCCAAA	120
Db	98	ACTTATATGAGCAATTTGGCGCCAGAACACACAGCTGAGCAAGAGGCCCTCCAAAGCCAAA	157
OY	121	GTTGATGCTGAGAACTTCTACTANTTCATTCCTGTACCTCATGCTGATGATTTGGAAATGTC	180
Db	158	GTTGATGCTGAGAACTTCTACTANTTCATTCCTGTACCTCATGCTGATGATTTGGAAATGTC	217
OY	181	TCTTTCATCATCAGTGGCCATCCCTGGTGGACACTGTGAAATCCGAAGAGAGCGGAACACTCC	240
Db	218	TCTTTCATCATCAGTGGCCATCCCTGGTGGACACTGTGAAATCCGAAGAGAGCGGAACACTCC	277
OY	241	AATGACCCCTACCCACAGTACATTTTATGAGGACTGGCAGGAAAAGTACAAAGGCCAAATTC	300
Db	278	AATGACCCCTACCCACAGTACATTTTATGAGGACTGGCAGGAAAAGTACAAAGGCCAAATTC	337
OY	301	TTGAAATAGAAAGAAATCGAAGGCCACCATCCATGAGAAACATTGGTGCGGCTGGGTTCAAA	360
Db	338	TTGAAATAGAAAGAAATCGAAGGCCACCATCCATGAGAAACATTGGTGCGGCTGGGTTCAAA	397
OY	361	ATGTCGCCC 369	
Db	398	ATGTCGCCC 406	

```
RESULT 6
AAK52645
ID AAK52645 standard; cDNA; 600 BP.
AC
XX AAK52645;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 2174.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation; ss.
XX
OS Homo sapiens.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US04098.
XX
PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
PA (HSE-) HISEQ INC.
XX
PI Tang YF, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QF, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX
DR WPI: 2001-476283/51.
DR P-PSDB; AAM79512.
XX
PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
XX
PS Claim 1: Page 4539-4540; 6221pp; English.
XX
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666
CC (AAM80020) are omitted as the relevant pages from the sequence listing
CC were missing at the time of publication.
XX
SQ Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;
Query Match 100.0%; Score 369; DB 22; Length 600;
Best Local Similarity 100.0%; Pred. No. 3e-101;
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGCTACTTATCCAAATTTACACAGACGCTGCGAAGAGTCTCCGAAGATTTTATT 60
DB 38 ATGCTACTTATCCAAATTTACACAGACGCTGCGAAGAGTCTCCGAAGATTTTATT 97
QY 61 ACTTATATGACAAATTTGGCGCCAGACACAGACGCTGAGCAGAGAGCCCTCCAAAGCCAAA 120
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DB 98 ACTTATATGACAAATTTGGCGCCAGACACAGACGCTGAGCAGAGAGCCCTCCAAAGCCAAA 157
QY 121 GTTGATGCTGAGAACTTCTACTATGTATCTGTACTGATCTGATGATGATGGAATGTTTC 180
DB 158 GTTGATGCTGAGAACTTCTACTATGTATCTGTACTGATCTGATGATGGAATGTTTC 217
QY 181 TCTTTCATCATGCTGAGCACTGCTGAGCACTGGAATCAAGAGAGGGAACATGCC 240
DB 218 TCTTTCATCATGCTGAGCACTGCTGAGCACTGGAATCAAGAGAGGGAACATGCC 277
QY 241 AATGACCCCTTACACACAGTATCTGTAGAGCACTGGCAGGAAAGTACAAGCCAAATC 300
DB 278 AATGACCCCTTACACACAGTATCTGTAGAGCACTGGCAGGAAAGTACAAGCCAAATC 337
QY 301 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAATGTTGCGGTGGTTCAA 360
DB 338 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAATGTTGCGGTGGTTCAA 397
QY 361 ATGTCCCCC 369
DB 398 ATGTCCCCC 406
RESULT 7
AAK51661
ID AAK51661 standard; cDNA; 655 BP.
XX
XX AAK51661;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 206.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation; ss.
XX
OS Homo sapiens.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US04098.
XX
PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
PA (HSE-) HISEQ INC.
XX
PI Tang YF, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QF, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX
DR WPI: 2001-476283/51.
DR P-PSDB; AAM78528.
XX
PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
XX
PS Claim 1: Page 1024; 6221pp; English.
XX
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
```

CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.

XX  
SO Sequence 655 BP; 196 A; 154 C; 146 G; 153 T; 6 other:

Query Match 100.0%; Score 369; DB 22; Length 655;  
Best Local Similarity 100.0%; Pred. No. 3.1e-101;

Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCTACTTATTCATTTACACAGACGCTGGAGAGCGTCCGAAAGATTATTTAT 60  
DB 93 ATGCTACTTATTCATTTACACAGACGCTGGAGAGCGTCCGAAAGATTATTTAT 152  
QY 61 ACTTATATGCAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 120  
DB 153 ACTTATATGCAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 212  
QY 121 GTTATATGCTGAACTTCTACTATATGTCATCTGTAAGTGTGATGTAATGTTTC 180  
DB 213 GTTATATGCTGAACTTCTACTATATGTCATCTGTAAGTGTGATGTAATGTTTC 272  
QY 181 TCTTTCATCATCTGCGCATCTGCTGTGAGCAGCTGTAATTCAGAGAGCCGAACTCC 240  
DB 273 TCTTTCATCATCTGCGCATCTGCTGTGAGCAGCTGTAATTCAGAGAGCCGAACTCC 332  
QY 241 AATGACCCCTACACAGATCATTTGTAGAGAGCTGCGAGAAAAGTACAAAGCCCAATC 300  
DB 333 AATGACCCCTACACAGATCATTTGTAGAGAGCTGCGAGAAAAGTACAAAGCCCAATC 392  
QY 301 TTGAATCTAGAGAAATGGAAGGCCACCATCATGAGAAATGTTGGCTGGTTCAAA 360  
DB 393 TTGAATCTAGAGAAATGGAAGGCCACCATCATGAGAAATGTTGGCTGGTTCAAA 452  
QY 361 ATGTCCCCC 369  
DB 453 ATGTCCCCC 461

RESULT 8

AA64071 ID AAC64071 standard; cDNA; 732 BP.

XX AAC64071:

XX 19-FEB-2001 (first entry)

DE Human potassium channel protein KCNE2 (MIRP1) cDNA, SEQ ID NO-1.

XX Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KM Mink-related; long QT syndrome; cardiac arrhythmia;  
KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KM HERG; ss.

XX Homo sapiens.

XX WO200063434-A1.

XX 26-OCT-2000.

XX 14-APR-2000; 2000WO-US10004.

XX 15-APR-1999; 99US-0129404.

PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYTA ) UNIV YALE.

PI Abbott GW, Seethi F, Splawski I, Keating MT, Goldstein SAN;

XX WPI; 2000-672747/65.

DR P-PSDB; AAB29585.

PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for

PS Claim 1; Page 118-119; 132pp; English.

CC The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-KR), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents cDNA encoding human KCNE2 (MIRP1).

XX  
SO Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other:

Query Match 100.0%; Score 369; DB 21; Length 732;  
Best Local Similarity 100.0%; Pred. No. 3.3e-101;

Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCTACTTATTCATTTACACAGACGCTGGAAGAGCTTCCGAAGATTATTTAT 60  
DB 74 ATGCTACTTATTCATTTACACAGACGCTGGAAGAGCTTCCGAAGATTATTTAT 133  
QY 61 ACTTATATGCAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 120  
DB 134 ACTTATATGCAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 193  
QY 121 GTTATATGCTGAACTTCTACTATATGTCATCTGTAAGTGTGATGTAATGTTTC 180  
DB 194 GTTATATGCTGAACTTCTACTATATGTCATCTGTAAGTGTGATGTAATGTTTC 253  
QY 181 TCTTTCATCATCTGCGCATCTGCTGTGAGCAGCTGTAATTCAGAGAGCCGAACTCC 240  
DB 254 TCTTTCATCATCTGCGCATCTGCTGTGAGCAGCTGTAATTCAGAGAGCCGAACTCC 313  
QY 241 AATGACCCCTACACAGATCATTTGTAGAGAGCTGCGAGAAAAGTACAAAGCCCAATC 300  
DB 314 AATGACCCCTACACAGATCATTTGTAGAGAGCTGCGAGAAAAGTACAAAGCCCAATC 373  
QY 301 TTGAATCTAGAGAAATGGAAGGCCACCATCATGAGAAATGTTGGCTGGTTCAAA 360  
DB 374 TTGAATCTAGAGAAATGGAAGGCCACCATCATGAGAAATGTTGGCTGGTTCAAA 433  
QY 361 ATGTCCCCC 369  
DB 434 ATGTCCCCC 442

RESULT 9

ABR6573 ID ABR6573 standard; DNA; 732 BP.

XX	AAK86573;
AC	
XX	24-SEP-2002 (first entry)
DT	
XX	
XX	cDNA encoding human ether-a-g-go related interacting protein MiRP1.
DE	
KW	Human; human ether-a-go-go related gene; HERG; KCRL1; MiRP1;
KM	long QT syndrome; LQT; single nucleotide polymorphism; cardiac arrhythmia;
KW	potassium channel; ss; gene.
XX	
OS	Homo sapiens.
XX	
FH	Key
FT	CDS
FT	Location/Qualifiers
FT	74..445
FT	/tag= a
FT	/product= "MiRP1"
XX	
PX	WO200242735-A2.
XX	
PD	30-MAY-2002.
XX	
PF	30-OCT-2001; 2001WO-US45644.
PR	30-OCT-2000; 2000US-244340P.
PA	(UYVA-) UNIV VANDERBILT.
PI	Balser JR, George AL, Roden DM;
PS	WPI: 2002-527650/56.
DR	P-PSDB; AAU59168.
XX	
PT	Identifying a potassium channel activity modulator for drug design,
PT	comprises contacting a compound with a potassium channel and rat
PT	cerebellar cDNA library (KCRL) polypeptide, and determining activity -
XX	
PS	Claim 17; Page 162-163; 164pp; English.
XX	
CC	The invention relates to identifying (M1) a compound that modulates
CC	biological activity of a potassium channel (PC), by contacting a
CC	compound with a structure comprising a PC polypeptide and a polypeptide
CC	cloned from a rat cerebellar cDNA library (KCRL), and determining the
CC	activity of the PC polypeptide in the presence and absence of the
CC	compound, where a difference in the activities indicates modulation of
CC	biological activity of PC. Also include are identifying (M2) a candidate
CC	compound that modulates the biological activity of a complex comprising a
CC	human ether-a-go-go-related gene (HERG) channel polypeptide and a KCRL
CC	polypeptide, identifying (M3) a candidate compound as a modulator of KCRL
CC	expression, modulating (M4) PC function in a subject, comprising
CC	administering to the subject a substance that provides expression of a
CC	KCRL-encoding nucleic acid molecule in a cell or tissue, where modulated
CC	PC function is desired, screening (M5) for susceptibility to a drug-
CC	induced cardiac arrhythmia in a subject, comprising obtaining a
CC	biological sample from the subject and detecting a polymorphism of a KCRL
CC	gene in the biological sample from the subject, where the presence of the
CC	polymorphism indicates the susceptibility of the subject to a
CC	drug-induced cardiac arrhythmia, an oligonucleotide pair, where a first
CC	oligonucleotide of the pair hybridises to a first portion of a KCRL gene
CC	which includes a polymorphism of the KCRL gene, and the second
CC	oligonucleotide of the pair hybridises to a second portion of the KCRL
CC	gene that is adjacent to the first portion and a set of antisense
CC	oligonucleotide primers, suitable for amplifying a portion of a KCRL gene
CC	which includes a polymorphism of the KCRL gene. (M1) is useful for
CC	identifying a compound that modulates biological activity of PC,
CC	especially HERG, for modulating PC function (ie modulating HERG
CC	activity) in a mammal, by preparing a composition comprising the
CC	compound and administering the composition. The compound is useful for
CC	treating or preventing long QT syndrome (LQT) and is useful in drug
CC	designing. The present sequence encodes a HERG interacting
CC	protein MiRP1 (not defined).
XX	
50	Sequence 732 BP: 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match	Similarity	Score	DB	Length
Local	100.0%	369	DB 24	732
Matches	369	Conservative	0	Mismatches
			0	Indels
			0	Gaps
				0
1	ATGCTACTTTATCCAAATTCACACAGACGGCTGGAAGAGCTCTCCGAAGATTTTAT	60		
74	ATGCTACTTTATCCAAATTCACACAGACGGCTGGAAGAGCTCTCCGAAGATTTTAT	133		
61	ACTTATATGAGCAATTTGGCGCCAGAACACACAGCTGAGCAAGAGGCTCCAGCCAAA	120		
134	ACTTATATGAGCAATTTGGCGCCAGAACACACAGCTGAGCAAGAGGCTCCAGCCAAA	193		
121	GTTGATGCTGAGCACTTCTACTATGTCATCCGTACCTCATGCTGATGATTTGGAAATGTC	180		
134	GTTGATGCTGAGCACTTCTACTATGTCATCCGTACCTCATGCTGATGATTTGGAAATGTC	253		
181	TCTTTTCATCATGCTGAGCCATCCTGGTGAGCACTGTGAATTCACAGAGGAGCACTCC	240		
254	TCTTTTCATCATGCTGAGCCATCCTGGTGAGCACTGTGAATTCACAGAGGAGCACTCC	313		
241	AATGACCCCTACACCACTAGTCTTTAGAGCACTGGCAGGAAAAGTACAGACCAATC	300		
314	AATGACCCCTACACCACTAGTCTTTAGAGCACTGGCAGGAAAAGTACAGACCAATC	373		
301	TTGAATCTGAGAAATCGAAGGCCACATCCATGAGAAATCTGGTGGCTGGTTCAAA	360		
374	TTGAATCTGAGAAATCGAAGGCCACATCCATGAGAAATCTGGTGGCTGGTTCAAA	433		
361	ATGTCCCCC 369			
434	ATGTCCCCC 442			
RESULT 10				
AAD35170				
AAD35170	standard; DNA; 732 BP.			
AAD35170;				
25-JUL-2002	(first entry)			
Human KCNE2 wild type DNA.				
Human: Min-K related ion channel protein; MiRP1; ion channel disorder;				
KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; gene; ds.				
Homo sapiens.				
Key	Location/Qualifiers			
CDS	74..445			
FT	/*tag= a			
FT	/product= "Human MiRP1 protein"			
WO200222875-A2.				
21-MAR-2002.				
11-SEP-2001; 2001WO-US28332.				
11-SEP-2000; 2000US-231571P.				
(UYYA ) UNIV YALE.				
Goldstein SAN.				
WPI: 2002-362360/39.				
P-PSDB; AAE22095.				
Novel gene encoding Min-K related ion channel protein subunit and				
polymorphisms in this gene associated with antibiotic-induced long QT				
syndrome -				

PS Claim 9; Page 43; 49pp; English.  
XX  
CC The present invention relates to novel KCNE2 genes encoding Mln-K related  
CC (MIRP) 1 ion channel proteins and polymorphisms in these genes that are  
CC associated with ion channel disorders including antibiotic-induced long  
CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
CC 57 or 116 of MIRP1 polypeptide or a mutation at a nucleotide position  
CC encoding the amino acid positions is useful for diagnosing the presence  
CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
CC are useful in the development of new drug therapies which selectively  
CC target one or more KCNE2 polymorphisms that are associated with cardiac  
CC arrhythmias. The present sequence is human KCNE2 wild type DNA.  
XX  
SQ Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other;

Query Match 100.0%; Score 369; DB 24; Length 732;  
Best Local Similarity 100.0%; Pred. No. 3.3e-101;  
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 ATGCTACTTATTCATTTTCACACAGAGCGTGGAGAGCGTCTCCGAAAGATTATTTAT 60  
Db 74 ATGCTACTTATTCATTTTCACACAGAGCGTGGAGAGCGTCTCCGAAAGATTATTTAT 133  
QY 61 ACTTATATGACAAATTGGCGCCAGACACAAACAGCTGAGCAAGAGCCCTCCAGCCAAA 120  
Db 134 ACTTATATGACAAATTGGCGCCAGACACAAACAGCTGAGCAAGAGCCCTCCAGCCAAA 193  
QY 121 GTTATGCTGGAACCTTCTACTATATCTCTGTAACCTCATGTGATGATTGAAATGTC 180  
Db 194 GTTATGCTGGAACCTTCTACTATATCTCTGTAACCTCATGTGATGATTGAAATGTC 253  
QY 181 TCTTTCATCATCTGGGCATCTGTGTGAGACCTGTGAATCCAGAGACGGAACTCC 240  
Db 254 TCTTTCATCATCTGGGCATCTGTGTGAGACCTGTGAATCCAGAGACGGAACTCC 313  
QY 241 AATGACCCCTACACACAGTATGTAGAGAGCTGGCAGGAAAAGTACAAAGCCCAATC 300  
Db 314 AATGACCCCTACACACAGTATGTAGAGAGCTGGCAGGAAAAGTACAAAGCCCAATC 373  
QY 301 TTGAATCTAGAAGATCGAAGGCCACATCATGAGAACATTGGTGGCGCTGGTTCAA 360  
Db 374 TTGAATCTAGAAGATCGAAGGCCACATCATGAGAACATTGGTGGCGCTGGTTCAA 433  
QY 361 ATGTCCCC 369  
Db 434 ATGTCCCC 442

RESULT 11  
AAC64083  
ID AAC64083 standard; DNA; 732 BP.  
XX  
AC AAC64083;  
XX  
DT 19-FEB-2001 (first entry)  
XX  
DE Human potassium channel protein KCNE2 (MIRP1) Q9E mutant DNA.  
XX  
KW Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KW Mink-related; long QT syndrome; cardiac arrhythmia;  
KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KW HERG; mutant; ds.  
XX  
OS Homo sapiens.  
OS Synthetic.  
OS  
PN WO200063434-A1.  
XX  
XX 26-OCT-2000.  
XX  
PF 14-APR-2000; 200DMO-US10004.  
XX

PR 15-APR-1999; 99US-0129404.  
XX  
PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYXA ) UNIV YALE.  
XX  
PI Abbott GW, Seati F, Splawski I, Keating MT, Goldstein SAN;  
XX WPI; 2000-672747/95.  
DR P-PSDB; AAB29593.  
XX  
PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
PT diagnosing and treating ion channel disorders, especially long QT  
PT syndrome -  
XX  
PS Claim 56; Page -; 132pp; English.

XX  
CC The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-KR), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (MIRP1) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.

CC Note: The present sequence is not shown in the specification, but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.  
XX

SQ Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other;

Query Match 99.6%; Score 367.4; DB 21; Length 732;  
Best Local Similarity 99.7%; Pred. No. 9.9e-101;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCTACTTATTCATTTTCACACAGAGCGTGGAGAGCGTCTCCGAAAGATTATTTAT 60  
Db 74 ATGCTACTTATTCATTTTCACACAGAGCGTGGAGAGCGTCTCCGAAAGATTATTTAT 133  
QY 61 ACTTATATGACAAATTGGCGCCAGACACAAACAGCTGAGCAAGAGCCCTCCAGCCAAA 120  
Db 134 ACTTATATGACAAATTGGCGCCAGACACAAACAGCTGAGCAAGAGCCCTCCAGCCAAA 193  
QY 121 GTTATGCTGGAACCTTCTACTATATCTCTGTAACCTCATGTGATGATTGAAATGTC 180  
Db 194 GTTATGCTGGAACCTTCTACTATATCTCTGTAACCTCATGTGATGATTGAAATGTC 253  
QY 181 TCTTTCATCATCTGGGCATCTGTGTGAGACCTGTGAATCCAGAGACGGAACTCC 240  
Db 254 TCTTTCATCATCTGGGCATCTGTGTGAGACCTGTGAATCCAGAGACGGAACTCC 313  
QY 241 AATGACCCCTACACACAGTATGTAGAGAGCTGGCAGGAAAAGTACAAAGCCCAATC 300  
Db 314 AATGACCCCTACACACAGTATGTAGAGAGCTGGCAGGAAAAGTACAAAGCCCAATC 373  
QY 301 TTGAATCTAGAAGATCGAAGGCCACATCATGAGAACATTGGTGGCGCTGGTTCAA 360  
Db 374 TTGAATCTAGAAGATCGAAGGCCACATCATGAGAACATTGGTGGCGCTGGTTCAA 433  
QY 361 ATGTCCCC 369

Db 434 ATGTCCCC 442

|||||

## RESULT 12

AAC64084

ID AAC64084 standard; DNA; 732 BP.

XX

AC AAC64084;

XX

DT 19-FEB-2001 (first entry)

XX

DE Human potassium channel protein KCNE2 (MIRP1) M54T mutant DNA.

XX

KW Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;

KM Mink-related; long QT syndrome; cardiac arrhythmia;

KW drug screening; knockout mouse; transgenic animal; ion channel disorder;

KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;

KW HERG; mutant; ds.

XX

OS Homo sapiens.

OS Synthetic.

XX

PN WO20063434-A1.

XX

PD 26-OCT-2000.

XX 14-APR-2000; 2000WO-US10004.

XX

PF 15-APR-1999; 9905-0129404.

XX

PA (UTAH ) UNIV UTAH RES FOUND.

XX (UYVA ) UNIV YALE.

XX

PI Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAM;

DR WPI; 2000-672747/65.

DR P-PSDB; AAB29594.

XX

PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for

PT diagnosing and treating ion channel disorders, especially long QT

PT syndrome -

XX

PS Claim 56; Page -; 132pp; English.

XX

CC The invention relates to novel ion channel proteins related to

CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of

CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,

CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,

CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,

CC respectively). The cDNAs encoding these proteins are given in AAC64071-

CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier

CC potassium channels (I<sub>KR</sub>), mutations in which are associated with long

CC QT syndrome. The invention also relates to methods of diagnosing long QT

CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a

CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic

CC nonhuman animals comprising a heterologous ion channel protein gene

CC of the invention, a transgenic animal comprising human KCNE2 and HERG

CC DNA, and methods of and screening drugs for treating long QT syndrome

CC using KCNE2 proteins (including mutants), nucleic acids encoding them

CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic

CC acids, and proteins may be used for diagnosing or treating ion channel

CC disorders, especially long QT syndrome. Transgenic animals comprising

CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.

CC The present sequence represents DNA encoding a mutant human KCNE2

CC (MIRP1) specifically claimed for use in diagnostic and drug screening

CC methods of the invention.

CC Note: The present sequence is not shown in the specification, but is

CC derived from the wild-type human KCNE2 cDNA sequence shown on page

XX 118-119.

SQ Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match 99.6%; Score 367.4; DB 21; Length 732;  
Best Local Similarity 99.7%; Pred. NO. 9.9e-101;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCTACTTTATCCAAATTTCCACAGACGCTGGAAGACGCTCTCCGAAGATTTTAT 60  
DB 74 ATGCTACTTTATCCAAATTTCCACAGACGCTGGAAGACGCTCTCCGAAGATTTTAT 133  
QY 61 ACTTATATGACAAATTTGGCCGAGAACACACACCTGAGCAAGAGCCCTCCAAACCA 120  
DB 134 ACTTATATGACAAATTTGGCCGAGAACACACACCTGAGCAAGAGCCCTCCAAACCA 193  
QY 121 GTTGATGCTGGAAGCTTCTATATGTCATCTGTACCTCATGAGTGAATGGAATGTC 180  
DB 134 GTTGATGCTGGAAGCTTCTATATGTCATCTGTACCTCATGAGTGAATGGAATGTC 253  
QY 181 TCTTTCATCATCTGTGGCCATCTCTGTGAGCACTGTGAATCCAAAGACGGAACATCC 240  
DB 254 TCTTTCATCATCTGTGGCCATCTCTGTGAGCACTGTGAATCCAAAGACGGAACATCC 313  
QY 241 AATGACCCCTTACACCACTATGATGAGGACTGGCAGAAAGTACAAAGCCAAATC 300  
DB 314 AATGACCCCTTACACCACTATGATGAGGACTGGCAGAAAGTACAAAGCCAAATC 373  
QY 301 TTGAATCTAGAAGATCGAAGCCACCATCCATGAGAAATTTGGTGGGCTGTCAAA 360  
DB 374 TTGAATCTAGAAGATCGAAGCCACCATCCATGAGAAATTTGGTGGGCTGTCAAA 433  
QY 361 ATGTCCCC 369  
DB 434 ATGTCCCC 442

## RESULT 13

AAC64085

ID AAC64085 standard; DNA; 732 BP.

XX

AC AAC64085;

XX

DT 19-FEB-2001 (first entry)

XX

DE Human potassium channel protein KCNE2 (MIRP1) 157T mutant DNA.

XX

KW Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;

KM Mink-related; long QT syndrome; cardiac arrhythmia;

KW drug screening; knockout mouse; transgenic animal; ion channel disorder;

KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;

KW HERG; mutant; ds.

XX

OS Homo sapiens.

OS Synthetic.

PN WO20063434-A1.

XX

PD 26-OCT-2000.

XX 14-APR-2000; 2000WO-US10004.

XX

PF 15-APR-1999; 9905-0129404.

XX

PA (UTAH ) UNIV UTAH RES FOUND.

XX (UYVA ) UNIV YALE.

XX

PI Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAM;

DR WPI; 2000-672747/65.

DR P-PSDB; AAB29595.

XX

PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for

PT diagnosing and treating ion channel disorders, especially long QT

PT syndrome -

XX

PS Claim 56; Page -; 132pp; English.

XX The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MiRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MiRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MiRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-Kr), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (MiRP1) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.  
CC Note: The present sequence is not shown in the specification, but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.  
XX  
XX  
SQ Sequence 732 BP; 221 A; 153 C; 157 G; 201 T; 0 other;

Query Match 99.6%; Score 367.4; DB 21; Length 732;  
Best Local Similarity 99.7%; Pred. No. 9.9e-101;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCTACTTATTCATTCACACAGACGCTGGAGAGCGTCTCCGAGAGATTTTAT 60  
DB 74 ATGCTACTTATTCATTCACACAGACGCTGGAGAGCGTCTCCGAGAGATTTTAT 133  
QY 61 ACTTATATGGACAATTTGGCGCCAGACACAAACAGCTGAGAGAGGCGCTCCAAAGCCAA 120  
DB 134 ACTTATATGGACAATTTGGCGCCAGACACAAACAGCTGAGAGAGGCGCTCCAAAGCCAA 193  
QY 121 GTTGATGCTGAGAACTTCTATATGATGATCTGATCTGATGATGATGATGATGATG 180  
DB 194 GTTGATGCTGAGAACTTCTATATGATGATCTGATCTGATGATGATGATGATGATG 253  
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DB 254 TCTTTCATCATCGTGGCCATCTGCTGAGACATGTGAAATCCAAAGAGCGGAACACTCC 313  
QY 241 AATGACCCCTACACACAGTACATTTGTAGAGACTGGCAGAAAAGTCAAGAGCCAAATC 300  
DB 314 AATGACCCCTACACACAGTACATTTGTAGAGACTGGCAGAAAAGTCAAGAGCCAAATC 373  
QY 301 TTGAATCTAGAGAAATGGAAGGCGCACCATCATGAGAAATTTGGTGGGCTGGTCAAA 360  
DB 374 TTGAATCTAGAGAAATGGAAGGCGCACCATCATGAGAAATTTGGTGGGCTGGTCAAA 433  
QY 361 ATGTCCCCC 369  
DB 434 ATGTCCCCC 442

RESULT 14  
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ID AAC64086 standard; DNA; 732 BP.  
XX  
AC AAC64086;  
XX  
DT 19-FEB-2001 (first entry)  
XX  
DE Human potassium channel protein KCNE2 (MiRP1) T8A mutant DNA.  
XX  
KW Human; KCNE2; MiRP1; potassium channel protein; KCNE1-related;

KW Mink-related; long QT syndrome; cardiac arrhythmia;  
KW drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KW fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KW HERG; mutant; ds.  
OS Homo sapiens.  
OS Synthetic.  
PN WO200063434-A1.  
XX  
PD 26-OCT-2000.  
XX  
PF 14-APR-2000; 2000MO-US10004.  
XX  
PR 15-APR-1999; 99US-0129404.  
XX  
PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UTAH ) UNIV YALE.  
XX  
PI Abbott GW, Seatl F, Splawski I, Keating MT, Goldstein SAN;  
DR F-PSDB; AAB29586.  
DR  
PT Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for  
PT diagnosing and treating ion channel disorders, especially long QT  
PT syndrome -  
XX  
XX  
PS Claim 56; Page -; 132pp; English.

CC The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MiRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MiRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MiRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-Kr), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (MiRP1) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.  
CC Note: The present sequence is not shown in the specification, but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.  
XX  
SQ Sequence 732 BP; 220 A; 152 C; 158 G; 202 T; 0 other;

Query Match 99.6%; Score 367.4; DB 21; Length 732;  
Best Local Similarity 99.7%; Pred. No. 9.9e-101;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCTACTTATTCATTCACACAGACGCTGGAGAGCGTCTCCGAGAGATTTTAT 60  
DB 74 ATGCTACTTATTCATTCACACAGACGCTGGAGAGCGTCTCCGAGAGATTTTAT 133  
QY 61 ACTTATATGGACAATTTGGCGCCAGACACAAACAGCTGAGAGAGGCGCTCCAAAGCCAA 120  
DB 134 ACTTATATGGACAATTTGGCGCCAGACACAAACAGCTGAGAGAGGCGCTCCAAAGCCAA 193  
QY 121 GTTGATGCTGAGAACTTCTATATGATGATCTGATCTGATGATGATGATGATGATG 180  
DB 194 GTTGATGCTGAGAACTTCTATATGATGATCTGATCTGATGATGATGATGATGATG 253



QY 181 TCTTTCATCATGTCGGCCATCCCTGCTGAGACACTGTGAATCCAAAGAGAGCCAACTCC 240  
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Db 254 TCTTTCATCATCTGCGCCATCCCTGCTGAGACACTGTGAATCCAAAGAGAGCCAACTCC 313  
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Db 374 TTGAATCTAGAAAGATCGAAGGCGCCATCCATGAGAAATTTGGTGGCTGGGTCAA 433  
QY 361 ATGTCCCCC 369  
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Db 434 ATGTCCCCC 442

RESULT 15  
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ID AAD35169 standard; DNA; 732 BP.  
XX  
AC AAD35169;  
XX  
DT 25-JUL-2002 (first entry)  
XX  
DE Human KCNE2 mutant DNA (C420T).  
XX  
KW Human; Min-K related ion channel protein; MiRP1; ion channel disorder;  
KM KCNE2; long QT syndrome; LQTS; cardiac arrhythmia; mutant; gene; SNP;  
KW single nucleotide polymorphism; ds.  
XX  
OS Homo sapiens.  
XX  
FH Key location/Qualifiers  
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FT FT /tag= a  
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FT /standard\_name= "Single nucleotide polymorphism (SNP)"  
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XX  
PD 21-MAR-2002.  
XX  
PF 11-SEP-2001; 2001WO-US28332.  
XX  
PR 11-SEP-2000; 2000US-231571P.  
XX  
PA (UYVA ) UNIV YALE.  
XX  
PI Goldstein SAN;  
XX  
DR WPI; 2002-362360/39.  
DR P-PSDB; AAE22094.  
XX  
PT Novel gene encoding Min-K related ion channel protein subunit and  
PT polymorphisms in this gene associated with antibiotic-induced long QT  
PT syndrome -  
XX  
PS Claim 1; Page 41-42; 49pp; English.  
XX  
CC The present invention relates to novel KCNE2 genes encoding Min-K related  
CC (MiRP) 1 ion channel proteins and polymorphisms in these genes that are  
CC associated with ion channel disorders including antibiotic-induced long  
CC QT syndrome (LQTS). Detecting a mutation at amino acid positions 8, 54,  
CC 57 or 116 of MiRP1 polypeptide or a mutation at a nucleotide position  
CC encoding the amino acid positions is useful for diagnosing the presence  
CC of a polymorphism that causes drug-induced LQTS. The diagnostic methods  
CC are useful in the development of new drug therapies which selectively  
CC target one or more KCNE2 polymorphisms that are associated with cardiac  
CC arrhythmias. The present sequence is human KCNE2 mutant DNA (C420T).

XX  
SQ Sequence 732 BP; 221 A; 151 C; 157 G; 203 T; 0 other;  
Query Match 99.6%; Score 367.4; DB 24; Length 732;  
Best Local Similarity 99.7%; Pred. No. 9.9e-101;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 ATGTCTACTTTCATTCATTTTCACAGAGCGCTGGAAAGAGCTTCCGAAGATTTTATTT 60  
Db 74 ATGTCTACTTTCATTCATTTTCACAGAGCGCTGGAAAGAGCTTCCGAAGATTTTATTT 133  
QY 61 ACTTATATGAGACAATTTGGCGCCAGAACACAAAGCTGAGCAAGAGGCCCTCCAAAGCCAAA 120  
Db 134 ACTTATATGAGACAATTTGGCGCCAGAACACAAAGCTGAGCAAGAGGCCCTCCAAAGCCAAA 193  
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Db 194 GTTGATGCTGAGAACTTTCTACTATGTCTCTGTACCTCATGTGATGATGATTTGAATGTTTC 253  
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QY 301 TTGAATCTAGAAAGATCGAAGGCGCCATCCATGAGAAATTTGGTGGCTGGGTCAA 360  
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QY 361 ATGTCCCCC 369  
Db 434 ATGTCCCCC 442

Search completed: May 21, 2003, 20:27:32  
Job time : 95.8202 secs



GenCore version 5.1.4.p5.4578  
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## OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 19:44:14 ; Search time 803.354 Seconds  
(without alignments)  
13367.622 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_442

Perfect score: 369

Sequence: 1 atgtctacttaccatc.....ctgggttcacaaatgcctccc 369

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapept 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: gb\_in:\*

4: gb\_om:\*

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6: gb\_pal:\*

7: gb\_ph:\*

8: gb\_pl:\*

9: gb\_pr:\*

10: gb\_ro:\*

11: gb\_scs:\*

12: gb\_sy:\*

13: gb\_un:\*

14: gb\_vl:\*

15: em\_ba:\*

16: em\_fun:\*

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24: em\_ph:\*

25: em\_pl:\*

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32: em\_hhg\_other:\*

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35: em\_hhg\_rtd:\*

36: em\_hhg\_mam:\*

37: em\_hhg\_vrt:\*

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40: em\_hhg\_mus:\*

41: em\_hhg\_other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

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2	369	100.0	732	9	AF071002
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6	369	100.0	100000	9	AP000167
7	369	100.0	100000	17	AP000120
8	369	100.0	340000	9	AF001719
9	367.4	99.6	732	6	AX406939
10	367.4	99.6	732	6	AX406943
11	367.4	99.6	732	6	AX406945
12	367.4	99.6	732	6	AX406947
13	265	71.8	1664	10	BC022699
14	263.4	71.4	372	10	AY050513
15	263.4	71.4	468	10	AF071003
16	261.8	70.9	144709	2	AC117904
17	186.6	50.6	225	4	AY079211
18	186.2	50.5	215	4	AF329636
19	178.2	48.3	228	4	AF387764
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21	55	14.9	750	10	GPI1SK
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23	53.2	14.4	390	10	AY050512
24	53.2	14.4	398	6	I40373
25	53.2	14.4	402	9	HUM1SKA
26	53.2	14.4	406	9	HUMCDPCA
27	53.2	14.4	436	9	HUM1SK
28	53.2	14.4	1703	6	AR119312
29	53.2	14.4	1703	6	AR164693
30	53.2	14.4	3173	9	BC036452
31	53.2	14.4	43126	9	AP000324
32	53.2	14.4	100000	9	AP000053
33	53.2	14.4	100000	9	AP000168
34	53.2	14.4	100000	17	AP000121
35	53.2	14.4	340000	9	AP001720
36	52.8	14.3	611	10	MMKCHA
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## ALIGNMENTS

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LOCUS

DEFINITION Sequence 3 from Patent WO0222875.

ACCESSION AX406941

VERSION AX406941.1 GI:21439816

KEYWORDS

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1

AUTHORS Goldstein,S.A.

TITLE Polymorphisms associated with cardiac arrhythmia

JOURNAL Patent: WO 0222875-A 3 21-MAR-2002;

Submitted (05-JUN-1998) Section of Developmental Biology and Biophysics, Department of Pediatrics and Boyer Center for Molecular

DOMENECH, A., ESTIVILL, X. and de la LUNA, S.  
Direct Submission

JOURNAL Submitted (01-SEP-2000) Medical and Molecular Genetics Center,  
Institut Recerca Oncologica, Avda. de Castelldefels Km 2.7,  
L'Hospitalet de Llobregat, Barcelona 08907, Spain

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BASE COUNT 247 a 172 c 189 g 200 t 1 others

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Query Match 100.0%; Score 369; DB 9; Length 809;  
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Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGTCTACTTTATCCAAATTCACACAGACGCTGGAAGCGTTCCGAAGATTTTATT 60  
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QY 61 ACTTATATGACAAATTTGGCGCCAGAACACACAGCTGACAGAGAGCCCTCCAGCCAA 120  
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QY 361 ATGTCCCC 369  
DB 501 ATGTCCCC 509

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DEFINITION clone:012C8, complete sequence.  
ACCESSION AP000320  
VERSION AP000320.1 GI:4835689  
KEYWORDS HTG.  
SOURCE Homo sapiens DNA, clone:Q12C8.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
AUTHORS Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Homo sapiens 24,608bp genomic DNA of 21q22.1  
JOURNAL Published Only in Database (1999)

REFERENCE 2 (bases 1 to 24608)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Direct Submission  
JOURNAL Submitted (13-MAY-1999) Masahira Hattori, The Institute of Physical  
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),  
Katsushika Univ., 1-15-1 Katsushika, Sagami-hara, Kanagawa 228-8555,  
Japan (E-mail:hattori@gsc.riken.go.jp,  
URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,  
Fax:81-42-778-9924)  
COMMENT The sequence is a part of the data (ACCESSION No. AP000165 -  
AP000173).  
The sequencing project is supported by Japan Science Technology  
Corporation (JST) and The Institute of Physical and Chemical  
Research (RIKEN).  
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Best Local Similarity 100.0%; Pred. No. 1.9e-100;  
Matches 369; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DEFINITION complete sequence.  
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VERSION AP000052.1 GI:3132362  
KEYWORDS HTG.  
SOURCE Homo sapiens DNA, clone:245P17-f4A4E\_2.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
AUTHORS I (bases 1 to 100000)  
TITLE Homo sapiens genomic DNA, chromosome 21q

JOURNAL  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
TITLE Direct Submission  
JOURNAL Submitted (11-MAY-1998) Masahira Hattori, Kitasato University,  
Department of Science, JST Sequencing Laboratory, Kitasato 1-15-1,  
Sagamihara 228, Japan (E-mail:hattori@engc.ims.u-tokyo.ac.jp,  
Tel:0427-78-9732, Fax:0427-78-9561)  
COMMENT This sequence is conducted by Kitasato University JST sequencing  
Laboratory as a JST sequencing team.  
JOURNAL Principal Investigator:Yoshiyuki Sakaki Ph.D.  
PHONE: +81-3-5449-5622, Fax: +81-3-5449-5445,  
sakaki@engc.ims.u-tokyo.ac.jp  
Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D. The  
sequence is submitted by Human Genome Sequencing in ALIS project of  
JST  
Japan Science and Technology Corporation (JST)  
5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0028 Japan  
For further information about this sequence, including its location  
and relationship to other sequences, please visit our sequence  
archive Web site (http://www.alls.tokyo.jst.go.jp/HGS/top.html)  
or send email to webmaster@www.alls.tokyo.jst.go.jp'.  
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QY 61 ACTTATATGACAAATTGGCCGCAACACAGAGCTGAGCAAGAGCCCTCCAGCCAA 120  
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QY 121 GTTGATGCTGAGAACTCTCTATGTCATCTGTACTCATGTGATGATTTGAATGTTTC 180  
DB 80338 GTTGATGCTGAGAACTCTCTATGTCATCTGTACTCATGTGATGATTTGAATGTTTC 80397  
QY 181 TCTTTCATCATGCTGGCCATCTGTGAGCACTGTGAATCCAAAGAGCGGAACACTCC 240  
DB 80398 TCTTTCATCATGCTGGCCATCTGTGAGCACTGTGAATCCAAAGAGCGGAACACTCC 80457  
QY 241 AATGACCCCTACACACAGTACATTTGTAGAGAGCTGGCAGGAAAAAGTACAGAGCCAAATC 300  
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QY 361 ATGTCCCCC 369  
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LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AML region,  
DEFINITION Clone B2344F14-f30E8, segment 3/9, complete sequence.  
ACCESSION AP000167  
VERSION AP000167.1 GI:4827132  
KEYWORDS HTG.

SOURCE Homo sapiens DNA.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
REFERENCE 1 (bases 1 to 100000)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Homo sapiens 890,291bp genomic DNA of 21q22.1 (REGION: D21S226-AML  
CLONE RANGE: B2344F14-150E8)  
JOURNAL Published Only in Database (1999)  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Direct Submission  
JOURNAL Submitted (10-MAY-1999) Masahira Hattori, The Institute of Physical  
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),  
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,  
Japan (E-mail:hattori@engc.riken.go.jp,  
URL:http://hgp.gsc.riken.go.jp, Tel:81-42-778-9923,  
Fax:81-42-778-9924)  
COMMENT E. coli transposon insertion: The present data does not contain E.  
coli transposon sequences which integrated in the  
original/previous sequences. We determined the boundary between  
the insertion and genomic sequences experimentally, removed the  
insertion sequences, reconstituted the present data. The sequencing  
project is supported by Japan Science Technology Corporation (JST)  
and The Institute of Physical and Chemical Research (RIKEN).  
FEATURES  
Location/Qualifiers  
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DB 72246 ACTTATATGACAAATTGGCCGCAACACAGAGCTGAGCAAGAGCCCTCCAGCCAA 72305  
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DB 72306 GTTGATGCTGAGAACTCTCTATGTCATCTGTACTCATGTGATGATTTGAATGTTTC 72365  
QY 181 TCTTTCATCATGCTGGCCATCTGTGAGCACTGTGAATCCAAAGAGCGGAACACTCC 240  
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QY 241 AATGACCCCTACACACAGTACATTTGTAGAGAGCTGGCAGGAAAAAGTACAGAGCCAAATC 300  
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QY 361 ATGTCCCCC 369  
DB 72546 ATGTCCCCC 72554  
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ID AP000120  
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AC Ap000120;  
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 SV Ap000120.1  
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 DT 04-MAY-1999 (Rel. 59, Created)  
 DT 26-SEP-1999 (Rel. 61, Last updated, Version 3)  
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 XX Homo sapiens genomic DNA of 21q22.1, GART and AML related, SLC5A3-f4A4  
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 XX HUG.  
 XX  
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 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;  
 CC Eutheria; Primates; Catarrhini; Homiidae; Homo.  
 XX  
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 RP 1-100000  
 RA Hirakawa M., Yamaguchi H., Imai K., Shimada J.;  
 RT  
 RL Submitted (15-APR-1999) to the EMBL/GenBank/DBJ databases.  
 RL Mika Hirakawa, Japan Science and Technology Corporation (JST), Advanced  
 RL Databases Department, 5-3, Yonbancho, Chiyoda-Ku, Tokyo 102-0081, Japan  
 RL (E-mail:mika@tokyo.jst.go.jp, URL:http://www-alls.tokyo.jst.go.jp/  
 RL Tel:81-3-5214-8491, Fax:81-3-5214-8470)  
 XX  
 XX [2]  
 RA Hattori M., Ishii K., Toyoda A., Shiba T., Sakaki Y.;  
 RL "Homo sapiens 817,199bp genomic DNA of 21q22.1 GART and AML region";  
 RL unpublished.  
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 CC This sequence is conducted by Kitasato University JST sequencing  
 CC Laboratory as a JST sequencing team.  
 CC Principal Investigator:Yoshiyuki Sakaki Ph.D.  
 CC Phone:+81-3-5449-5622, Fax:+81-3-5449-5445,  
 CC sakaki@ngc.ims.u-tokyo.ac.jp  
 CC Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D.  
 CC The sequence is submitted by Human Genome Sequencing in ALIS  
 CC project of JST.  
 CC Japan Science and Technology Corporation (JST)  
 CC 5-3, Yonbancho, Chiyoda-Ku, Tokyo 102-0081 Japan  
 CC For further information about this sequence, including its  
 CC location and relationship to other sequences, please visit our  
 CC sequence archive Web site (http://www-alls.tokyo.jst.go.jp/HGS/)  
 CC or send email to webmaster@www-alls.tokyo.jst.go.jp  
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 FT /sequenced\_mol="DNA"  
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 FT /map="21q22.1"  
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 FT STS  
 FT 43934..44157  
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 FT /note="AFM086y9:Genethon Marker:The location is between  
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 DEFINITION Homo sapiens genomic DNA, chromosome 21q, section 63/105.  
 ACCESSION AP001719 AL163264 BA000005  
 VERSION AP001719.1 GI:7768719  
 KEYWORDS  
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 ORGANISM  
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 1 (sites)  
 Hattori M., Fujiyama A., Taylor T.D., Watanabe H., Yada T.,  
 Park H.S., Toyoda A., Ishii K., Tokoi Y., Choi D.K., Soeda E.,  
 Ohki M., Takagi T., Sakaki Y., Taudien S., Blechschmidt K.,  
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 Antonarakis S.E., Minoshima S., Shimizu N., Nordisick G.,  
 Hornischer K., Barand P., Scharfe M., Schoen O., Desario A.,  
 Reichelt J., Kauer G., Bloeker H., Ramser J., Beck A., Klages S.,  
 Hennig S., Riesenmann L., Dagand E., Wehmeyer S., Borzym K.,  
 Gardiner K., Nizetic D., Francis F., Lehrach H., Reinhardt R. and  
 Vaspo M.L.  
 The DNA sequence of human chromosome 21  
 Nature 405 (6784), 311-319 (2000)  
 2 (bases 1 to 340000)  
 Hattori M., Fujiyama A., Taylor T.D., Watanabe H., Yada T.,  
 Park H.S., Toyoda A., Ishii K., Tokoi Y., Choi D.K., Soeda E.,  
 Ohki M., Takagi T., Sakaki Y., Taudien S., Blechschmidt K.,  
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 Patterson D., Reichwald K., Rump A., Schillhabel M., Schudy A.,  
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 Antonarakis S.E., Minoshima S., Shimizu N., Nordisick G.,  
 Hornischer K., Barand P., Scharfe M., Schoen O., Desario A.,  
 Reichelt J., Kauer G., Bloeker H., Ramser J., Beck A., Klages S.,  
 Hennig S., Riesenmann L., Dagand E., Wehmeyer S., Borzym K.,  
 Gardiner K., Nizetic D., Francis F., Lehrach H., Reinhardt R. and  
 Vaspo M.L.  
 Direct Submission  
 Submitted (10-APR-2000) The Chromosome 21 Mapping and Sequencing  
 Consortium \* RIKEN Genomic Sciences Center, Human Genome Research  
 Group \* Institute of Molecular Biotechnology, Genome Analysis \*  
 Keio University School of Medicine, Dept. of Molecular Biology \*  
 GNF, Dept. of Genome Analysis \* Max-Planck Institute for Molecular





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DEFINITION	Sequence 5 from Patent WO222875.							
ACCESSION	AX406943							
VERSION	AX406943.1	GI:21439818						
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SOURCE	human.							
ORGANISM	Homo sapiens							
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;							
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.							
TITLE	Goldstein, S. A.							
JOURNAL	Polymorphisms associated with cardiac arrhythmia							
FEATURES	Patent: WO 0222875-A 5 21-MAR-2002;							
SOURCE	YALE UNIVERSITY (US)							
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Best Local Similarity 99.7%: Pred. No. 2.6e-100;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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ACCESSION AX406945  
VERSION AX406945.1 GI:21439820  
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SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 Goldstein, S.A.  
AUTHORS Polymorphisms associated with cardiac arrhythmia  
TITLE Patent: WO 0222875-A 7 21-MAR-2002;  
JOURNAL VALE UNIVERSITY (US)  
FEATURES location/Qualifiers  
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variation 243  
/note="The drug associated here was oxalotomide."

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ACCESSION AX406947  
VERSION AX406947.1 GI:21439822  
KEYWORDS  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 Goldstein, S.A.  
AUTHORS Polymorphisms associated with cardiac arrhythmia  
TITLE Patent: WO 0222875-A 9 21-MAR-2002;  
JOURNAL VALE UNIVERSITY (US)  
FEATURES location/Qualifiers  
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BASE COUNT 220 a 152 c 158 g 202 t  
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Best Local Similarity 99.7%: Pred. No. 2.6e-100;  
Matches 368; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 74 ATGCTACTTTATTCACATTTTCACACAGACGCTGGAGAGCTCTCCGAAAGATTTTATT 133
OY 61 ACTTATATGACAAATTGGCGCCAGACAAACAGCTGAGCAAGAGCCCTCCAAAGCCAAA 120
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OY 121 GTTGATGCTGAAACTTCTACTATGTCATCCTGTACTCATGTGATGATTTGAATGTTTC 180
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 DEFINITION AF071003  
 ACCESSION AF071003.1 GI:4704424  
 VERSION  
 KEYWORDS  
 SOURCE Rattus norvegicus.  
 ORGANISM Rattus norvegicus.  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.  
 1 (bases 1 to 468)  
 Abbott, G.W., Sesti, F., Splawski, I., Buck, M.E., Lehmann, M.H.,  
 Timothy, K.W., Keating, M.T. and Goldstein, S.A.  
 MIRP1 forms IKR potassium channels with HERG and is associated with  
 cardiac arrhythmia  
 Cell 97 (2), 175-187 (1999)

REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL  
 MEDLINE  
 PUBMED  
 REFERENCE  
 AUTHORS  
 TITLE  
 JOURNAL

FEATURES  
 source  
 1..468  
 Location/Qualifiers  
 /organism="Rattus norvegicus"  
 /strain="Sprague-Dawley"  
 /db\_xref="taxon:10116"  
 /sex="male"  
 /tissue\_type="heart"  
 /dev\_stage="10-12 weeks"  
 35..406  
 /note="potassium channel subunit; MIRP1"  
 /codon\_start=1  
 /product="mink-related peptide 1"

CDS

/protein\_id="AAD28087.1"  
 /db\_xref="GI:4704425"  
 /translation="MTTLANLTOTLEDARKKVEITYMDSMRNTTAEDQALARVDAE  
 NEFYVLYLMWIMGMEAFIVVALIVSTVASKRREHSODPIHQIYIEDNQKRRSDILH  
 LEDSKATHENIGATCFIVSP"  
 BASE COUNT 118 a 126 c 111 g 93 t  
 ORIGIN

Query Match 71.4%; Score 263.4; DB 10; Length 468;  
 Best Local Similarity 82.1%; Pred. No. 8.3e-69;  
 Matches 303; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

```

OY 1 ATGCTACTTATTCATTTTCACACAGAGCGTGGAAAGCGTCTCCGAGATTTTAT 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 35 ATGACCACTTTAGCCACTTACGACGACCGCTGGAGTGCCCTTCAAAAAGTTTCAT 94
OY 61 ACTTATATGACAAATGGCGCCACACACACAGCTGAGAGCGCTCCAGCCAAA 120
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 95 ACTTATATGACAGCTGGAGAGAGACACACAGCGGAAACACAGCGCTCCAGGCCAGA 154
OY 121 GTTGATGCTGAAACTTCTACTATGTCATCCTGTACTCATGTGATGATTTGAATGTTTC 180
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 121 GTTGATGCTGAAACTTCTACTATGTCATCCTGTACTCATGTGATGATTTGAATGTTTC 180
OY 155 GTTGATGCTGAAACTTCTACTATGTCATCCTGTACTCATGTGATGATTTGAATGTTTC 214
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 181 TCTTTCATCATGTCGTCATCTGTGTGACATGTAATCCAGACAGCGGAACTCC 240
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 215 GCCTTCATCTGTGTGTCATCTGTGTGACAGCGTGAAGTCGAAGCGGGGAGCACTCC 274
OY 241 AATGACCCCTACACAGTACATTTGTAGAGACTGGCAGAAAGTACAAAGCCAAATC 300
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 275 CAGGACCGGTACACAGTACATTTGTAGAGACTGGCAGAAAGTATAGAGATCAGATC 334
OY 301 TTGAATCTAGAAAGATGGAAGGCCACATCCATGAGAAACATTTGGTGGGCTTCAAA 360
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 335 TTGATCTGGAAGACTCCAAAGCCACATCCATGAGAAACCTGGGGGAGCGGTTTACA 394
OY 361 ATGTCCCC 369
    ||| |||
Db 395 GTGTCAACC 403
  
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Search completed: May 21, 2003, 21:21:35  
 Job time : 1122.35 secs

Gencore version 5.1.4 p5.4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 15, 2003, 14:21:24 ; Search time 44 Seconds

(without alignments)  
268.739 Million cell updates/sec

Title: US-09-550-163-2

Perfect score: 632  
Sequence: 1 MSTLSNFTQTLIEDVRRIRFI.....EESKATIHENIGAGFKMSP 123

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :  
1: p1r1:\*  
2: p1r2:\*  
3: p1r3:\*  
4: p1r4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	132.5	21.0	129	1 A33447	potassium channel-
2	128.5	20.3	129	2 S17307	potassium channel
3	128	20.3	130	2 A35633	potassium channel
4	126.5	20.0	125	2 A49392	min K potassium ch
5	126.5	20.0	125	2 I48146	potassium channel-
6	84.5	13.4	439	2 D64510	hypothetical prote
7	76.5	12.1	374	2 T15940	hypothetical prote
8	76.5	12.1	557	2 B47704	probable sensory t
9	75.5	11.9	367	2 E81065	proteinase, probab
10	75.5	11.9	3839	2 T49799	related to TOM1 pr
11	75	11.9	1003	2 AH2335	toxin secretion AB
12	74.5	11.8	1042	2 H70203	isoletucine-tRNA Ii
13	74	11.7	938	2 T41932	hypothetical prote
14	73.5	11.6	171	2 T41924	hypothetical prote
15	73.5	11.6	497	2 S59103	NADH2 dehydrogenas
16	73.5	11.6	652	2 T41162	hypothetical prote
17	73	11.6	400	2 B64071	lysine-specific
18	73	11.6	591	2 AD2148	two-component sens
19	73	11.6	1224	2 E71611	hypothetical prote
20	72.5	11.5	371	2 D89995	accessory gene reg
21	71	11.2	174	2 AG2234	hypothetical prote
22	71	11.2	291	2 A36051	H+/K+-exchanging A
23	70.5	11.2	451	2 F75131	valyl-tRNA synthet
24	70.5	11.2	874	2 E82913	hypothetical prote
25	70	11.1	522	2 C90073	hypothetical prote
26	70	11.1	947	2 G70657	probable adl - Myc
27	70	11.1	1357	2 T16860	hypothetical prote
28	69.5	11.0	554	2 T27878	hypothetical prote
29	69.5	11.0	946	2 S48255	probable membrane

30	69.5	11.0	1154	2 T39663	paired amphipathic
31	69.5	11.0	3973	2 B71612	hypothetical prote
32	69	10.9	282	1 E69906	conserved hypotet
33	69	10.9	548	2 E89910	glycine betaine tr
34	69	10.9	773	2 D90099	hypothetical prote
35	68.5	10.8	350	2 B81803	probable secreted
36	68.5	10.8	655	2 H96692	probable receptor
37	68.5	10.8	696	2 C85047	probable transposo
38	68.5	10.8	701	2 S35313	TIP1 protein - yea
39	68.5	10.8	1007	2 S48535	rno-type GTPase-ac
40	68.5	10.8	1911	2 T43048	calcium channel al
41	68	10.8	303	2 T15694	hypothetical prote
42	68	10.8	378	2 G97301	uncharacterized co
43	68	10.8	435	2 B97516	hypothetical prote
44	68	10.8	435	2 A12734	ABC transporter, m
45	68	10.8	464	2 T20238	hypothetical prote

#### ALIGNMENTS

```

RESULT 1
A32447
potassium channel-activating protein - human
N:Altermate names: delayed rectifier potassium channel
C:Species: Homo sapiens (man)
C>Date: 28-Jan-2000 #sequence_revision 28-Jan-2000 #text_change 21-Jul-2000
C:Accession: A32447; I53911
R:Murai, T.; Kakiyama, A.; Takumi, T.; Ohkubo, H.; Nakanishi, S.
Biochem. Biophys. Res. Commun. 161, 176-181, 1989
A>Title: Molecular cloning and sequence analysis of human genomic DNA encoding a nove
A:Reference number: A32447; MUID:89273632; PMID:2730656
A:Accession: A32447
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-129 <MUR>
A:Cross-References: GB:M26685; NID:g186569; PIDN:AAA36129.1; PID:g386838
R:Iral, L.P.; Deng, C.L.; Moss, A.J.; Kass, R.S.; Liang, C.S.
Gene 151, 339-340, 1994
A>Title: Polymorphism of the gene encoding a human minimal potassium ion channel (min
A:Reference number: I53911; MUID:95129890; PMID:7828904
A:Accession: I53911
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-37, G' 39-129 <LA>
A:Cross-References: GB:I33815; NID:g603450; PIDN:AAA63905.1; PID:g603451
C:Genetics:
A:Gene: GDB:KCNEL; ISK
A:Cross-References: GDB:127909; OMIM:176261
A:Map position: 21q22.1-21q22.2
C:Superfamily: human potassium channel-activating protein
C:Keywords: phosphoprotein; transmembrane protein
F:44-06/Domain: transmembrane #status predicted <TM>

Query Match 21.0%; Score 132.5; DB 1; Length 129;
Best Local Similarity 45.1%; Pred. No. 3.6e-06;
Matches 23; Conservative 15; Mismatches 12; Indels 1; Gaps 1;

QY 51 LILVMWIGMSFIIIVAILVSTYKSRKRRHSNDPIQYVED-WQKYSQI 100
DB 45 LVLVWLIGFEGFTLIMLSYRSKRLKHSNDPFVYIESDAMQKDKAVY 95

RESULT 2
S17307
potassium channel protein - mouse
C:Species: Mus musculus (house mouse)
C>Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 10-Dec-1999
C:Accession: S17307; S21135
R:Honore, E.; Attali, B.; Romey, G.; Heurteaux, C.; Ricard, P.; Lesage, F.; Lazdunski
EMBO J. 10, 2805-2811, 1991
A>Title: Cloning, expression, pharmacology and regulation of a delayed rectifier K+
A:Reference number: S17307; MUID:92007723; PMID:1655403

```



C:Accession: D64510  
R:Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann, R.D.; Sutton, G.G.; Blake,  
Reich, C.I.; Overbeek, R.; Kirkness, E.F.; Weinstock, K.G.; Merrick, J.M.; Glodok, A.  
rson, J.D.; Sadow, P.W.; Hanna, M.C.; Cotton, M.D.; Roberts, K.M.; Hurst, M.A.  
Science 273, 1058-1073, 1996

A:Authors: Kaine, B.P.; Borodovsky, M.; Klek, H.P.; Fraser, C.M.; Smith, H.O.; Woese, C.  
A:Title: Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii  
A:Reference complete: Ab4300; MUID:9633799; PMID:868087

A>Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Accession: D64510

A:Molecule type: DNA  
A:Residues: 1-439 <BUL>

A:Cross-references: GB:L77118; NID:g1500644; TIGR:MJECU4; PIDN:AAC37078.1; PID:g1500653

C:Genetics:  
A:Map position: EC1FOR1934-3253

A:Genome: plasmid  
A>Note: this stable 58-kilobase pair plasmid is also designated ECL (large extrachromosomal)  
C:Superfamily: Methanobacterium thermoautotrophicum conserved hypothetical protein MTH19

Query Match 13.4%; Score 84.5; DB 2; Length 439;  
Best Local Similarity 23.3%; Pred. No. 0.92;  
Matches 24; Conservative 19; Mismatches 31; Indels 29; Gaps 3;

OY 33 EQEALQAKVDAENFYIYLTMWIMGFSTIVAILNVTSKRRH----- 79  
| : | | : | : | : | : | : | : | : | :  
Db 8 ELKMINKEDSNNEEFVIYGRRIKTKLAKSV-----ENNREHIYYLAEGDNLKH 60  
| : | | : | : | : | : | : | : | : | :  
OY 80 -----SNDPYHOTIVEDMOEKY---KSQILNLEESKATIH 113  
| : | | : | : | : | : | : | : | : | :  
Db 61 EKRYASKEPTIEYAKDEWEAFYNFLDKDIIIDEFPALKIN 103

```

RESULT 7
T15940
hypothetical protein F01E11.4 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 20-Sep-1999
R:Accession: T15940
R:Gelsel, C.
submitted to the EMBL Data Library, December 1995
A:Description: The sequence of C. elegans cosmid F01E11.
A:Reference number: Z18434
A:Accession: T15940
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-374 <GENI>
A:Cross-references: EMBL:U04283Z; NID:g1125742; PID:g1125746; PIDD:AAA83573.1; CESP:F01E11.4
A:Gene: CESP.F01E11.4
A:Introns: 76/1; 104/3; 193/3; 229/3; 244/1; 256/2; 282/1; 328/2

Query Match      12.1%; Score 76.5; DB 2; Length 374;
Best Local Similarity 19.7%; Pred. No. 4.9;
Matches 23; Conservative 20; Mismatches 41; Indels 57; Gaps 4;

QY    31 TAEEALQAKDAENFFVYLILMWIMGFSLIAI-----67
       | : | | | | | | | | | | | | | | | |
Db    117 TLREVSFLASVIGNLIYYCPYLILVLGLSFWMTVPFFESILLNDKIMITHPATIFF 176
       + : + + + + + + + + + + + + + + + +

QY    68 -----LVSTVKSKRRH-----SNDEHYOIVEDWQEKYS---QIL 101
       : : : : : : : : : : : : : : : :
Db    177 ATICAMTAILGYDINSIHSSSDMEHWMKHSIKEDYGNTAPAHIMEWNKAHQFCGCVR 236
       ||| : ||| : ||| : ||| : ||| : ||| :

QY    102 NLEE-----SKATIHENIGANGFK 120
       || : || : || : || : || : || : ||
Db    237 NLTPDEVNSCCASCATMHERECVAPEK 263

RESULT 8
B47704 .
probable sensory transducer protein - Clostridium thermocellum
C:Species: Clostridium thermocellum
C:Date: 21-Jan-1994 #sequence_revision 18-Nov-1994 #text_change 15-Oct-1999

```

```

C:Accession: B47704
R:Hailewood, G.P.; Davidson, K.; Laurie, J.I.; Huskisson, N.S.; Gilbert, H.J.
J. Gen. Microbiol. 139, 307-316, 1993
A:Title: Gene sequence and properties of Celli, a family E endoglucanase from Clostridi
A:Reference number: A47704; MUID:93171873; PMID:8436946
A:Accession: B47704
A:Status: Preliminary; not compared with conceptual translation
A:Molecule type: DNA
A:Residues: 1-557 <HAZ>
A:Cross-references: CB:L04736; NID:q144912; PIND:AAA20891.1; PID:q144913
A:Note: sequence extracted from NCBI backbone (NCBI:P125641)
C:Superfamily: methyl-accepting chemotaxis protein
C:Keywords: transmembrane protein

Query Match      12.1%; Score 76.5; DB 2; Length 557;
Best Local Similarity 23.2%; Pred. No. 7.6;
Matches 29; Conservative 24; Mismatches 47; Indels 25; Gaps 5;

Oy 1 MSTLSNFGOTLEDVRRFRFFITYMDMWRONTAEQALQAKDAENFYVILYLWVIGMF 60
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 84 LSDIDDIDGTDLNDYF-EAFVEY-----NTTAAKEKVDENKQVASTASTVAIVLEGIIL 136
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Oy 61 SFPIIAIVLSFYKSRREHSNDPYHOYL-----VE-DMQEKYSQILNEESKAT 109
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 137 IATAIGVTSIRLTK-----PIQGVYEAADRILAGVEYDVAKETPDEIGKLAESPKR 189
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Oy 110 IHENI 114
      : : : :
Db 130 MTENI 194

```

[illegible]









GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:15:00 : Search time 1800 Seconds

(without alignments)  
4324.818 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_445

Perfect score: 372  
Sequence: 1 atgtctactttatcctaatt.....ggtcaanaatgtccctctga 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0', Gapext 1.0

Searched: 1797656 segs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :  
1: GenEmbl:  
2: gb\_ba:  
3: gb\_hg:  
4: gb\_in:  
5: gb\_ov:  
6: gb\_pat:  
7: gb\_ph:  
8: gb\_pl:  
9: gb\_pr:  
10: gb\_ro:  
11: gb\_sts:  
12: gb\_sy:  
13: gb\_un:  
14: gb\_vl:  
15: em\_ba:  
16: em\_fun:  
17: em\_hum:  
18: em\_in:  
19: em\_inu:  
20: em\_om:  
21: em\_or:  
22: em\_ov:  
23: em\_pat:  
24: em\_ph:  
25: em\_pl:  
26: em\_ro:  
27: em\_sts:  
28: em\_un:  
29: em\_vl:  
30: em\_htg\_hum:  
31: em\_htg\_inv:  
32: em\_htg\_other:  
33: em\_htg\_inv:

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Query Match	Length	DB ID	Description
-----				

1	372	100.0	732	9	AF071002
2	372	100.0	809	9	AF302095
3	372	100.0	24608	9	AP000320
4	372	100.0	100000	9	AP000052
5	372	100.0	100000	9	AP000120
6	372	100.0	100000	9	AP000167
7	372	100.0	340000	9	AP001719
8	268	72.0	1664	10	BC022699
9	266.4	71.6	372	10	AY050513
10	266.4	71.6	468	10	AF071003
11	186.2	50.1	215	4	AF329636
12	178.2	47.9	528	4	AF387764
13	56	15.1	234	4	RABIPCS
14	55	14.8	750	10	GPITRK
15	53.2	14.3	390	10	AF135188
16	53.2	14.3	390	10	AY050512
17	53.2	14.3	398	6	I40373
18	53.2	14.3	408	9	HUMISKA
19	53.2	14.3	408	9	HUMCDRPCA
20	53.2	14.3	436	9	HUMISK
21	53.2	14.3	1703	6	AR119312
22	53.2	14.3	1703	6	AR164693
23	53.2	14.3	43126	9	AP000324
24	53.2	14.3	100000	9	AP000053
25	53.2	14.3	100000	9	AP000121
26	53.2	14.3	100000	9	AP000168
27	53.2	14.3	340000	9	AP001720
28	52.8	14.2	611	10	MMKCHA
29	52.6	14.1	393	4	AB032575
30	51	13.7	471	10	RATDTRCA
31	51	13.7	585	10	RATPCPA
32	50.4	13.5	390	4	MSU62404
33	44.6	12.0	422	12	SYNMINK
34	41	11.0	6408	6	AX346023
35	39	10.5	161482	2	AP004643
36	38.6	10.4	4881	3	DSPTERH2
37	38.2	10.3	166856	2	AC011944
38	38.2	10.3	177045	2	AC015970
39	38.2	10.3	179522	2	AC104231
40	37.2	10.0	346294	1	AP002999
41	36.6	9.8	2652	6	A75965
42	36.6	9.8	2652	6	160522
43	36.6	9.8	2861	9	S62035
44	36.6	9.8	11488	1	AE005321
45	36.6	9.8	11749	1	AE000212

## ALIGNMENTS

RESULT 1  
AF071002 LOCUS Homo sapiens milk-related peptide 1 mRNA, complete cds.  
DEFINITION AF071002.1 GI:4704422  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
human.  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE  
1 (bases 1 to 732)  
Abbott,G.W., Sesti,F., Splawski,I., Buck,M.E., Lehmann,M.H.,  
Timothy,K.W., Keating,M.T. and Goldstein,S.A.  
MIRP forms IKR potassium channels with HERG and is associated with  
cardiac arrhythmia  
Cell 97 (2), 175-187 (1999)  
JOURNAL MEDLINE  
99235979  
TITLES  
2 (bases 1 to 732)  
Abbott,G.W., Sesti,F., Buck,M.E. and Goldstein,S.A.N.  
Direct Submission  
Submitted (05-JUN-1998) Section of Developmental Biology and  
Biophysics, Department of Pediatrics and Boyer Center for Molecular

Medicine, Yale University School of Medicine, 295 Congress Avenue,  
New Haven, CT 06536, USA  
Location/Qualifiers

FEATURES  
source

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/tissue="heart"  
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/db\_xref="GI:4704423"  
/translation="MSTLSNFTQTLDEVFRRIFITYMDNMRONTAEQALQKVD  
NFYVILYLMVMHGMFSFIVAILVSTVSKRREHSNDPHYQIVEDMQEKYSQILN  
LEESKATIHENIGAGFRKSP"

## CDS

BASE COUNT 221 a 152 c 157 g 202 t  
ORIGIN

## Query Match

100.0%; Score 372; DB 9; Length 732;

Best Local Similarity 100.0%; Pred. No. 1.3e-100;  
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCTCACTTATTCACACAGACGCTGGAGACGCTCCGAGAGATTTTATT 60  
|||||  
DB 74 ATGCTCACTTATTCACACAGACGCTGGAGACGCTCCGAGAGATTTTATT 133  
|||||  
QY 61 ACTATATGACATTTGGCCGCAACAAACAGCTGAGCAAGAGGCCCCCAAGCCAAA 120  
|||||  
DB 134 ACTATATGACATTTGGCCGCAACAAACAGCTGAGCAAGAGGCCCCCAAGCCAAA 193  
|||||  
QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCTGACCTCATGATGATTTGAATGTTTC 180  
|||||  
DB 194 GTTGATGCTGAGAACTTCTACTATGTCATCTGACCTCATGATGATTTGAATGTTTC 253  
|||||  
QY 181 TCTTTATCATCTGCGGCATCTGCTGAGCACTGTGAATCCAAAGAGGGAACACTCC 240  
|||||  
DB 254 TCTTTATCATCTGCGGCATCTGCTGAGCACTGTGAATCCAAAGAGGGAACACTCC 313  
|||||  
QY 241 AATGACCCCTACACACAGTATGTAGAGACTGGCAGGAAAAGTCAAGAGCCAAATC 300  
|||||  
DB 314 AATGACCCCTACACACAGTATGTAGAGACTGGCAGGAAAAGTCAAGAGCCAAATC 373  
|||||  
QY 301 TTGAATCTAGAAAGATGGAAGGCCACCATCCATGAGAACATTGGTGGCTGGGTTCAAA 360  
|||||  
DB 374 TTGAATCTAGAAAGATGGAAGGCCACCATCCATGAGAACATTGGTGGCTGGGTTCAAA 433  
|||||  
QY 361 ATGTCCCCCTGA 372  
|||||  
DB 434 ATGTCCCCCTGA 445  
|||||

RESULT 2  
AF302095 809 bp mRNA linear PRI 14-SEP-2000  
LOCUS

DEFINITION Homo sapiens voltage-gated K+ channel subunit MIRP1 (KCNE2) mRNA,  
complete cds.

ACCESSION AF302095  
VERSION AF302095.1 GI:10121887

## KEYWORDS

## SOURCE

## ORGANISM

human.  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 809)  
AUTHORS Domenech,A., Estivill,X. and de la Luna,S.  
JOURNAL Cloning of human MIRP1 cDNA  
REFERENCE 2 (bases 1 to 809)  
AUTHORS Domenech,A., Estivill,X. and de la Luna,S.  
JOURNAL Direct Submision

JOURNAL Submitted (01-SEP-2000) Medical and Molecular Genetics Center,  
Institut Recerca Oncologica, Avia. de Castelldefels Km 2,7,  
L'Hospitalet de l'lobregat, Barcelona 08907, Spain  
Location/Qualifiers

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## CDS

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DB 141 ATGCTCACTTATTCACACAGACGCTGGAGACGCTCCGAGAGATTTTATT 200  
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QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCTGACCTCATGATGATTTGAATGTTTC 180  
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DB 501 ATGTCCCCCTGA 512  
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RESULT 3  
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LOCUS

DEFINITION Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AMU region,  
clone:Q12C8, complete sequence.

ACCESSION AP000320  
VERSION AP000320.1 GI:4835689

## KEYWORDS

## SOURCE

## ORGANISM

human.  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE 1 (bases 1 to 24608)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
JOURNAL Published Only in Database (1999) in press

REFERENCE 2 (bases 1 to 24608)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
 Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (13-MAR-1999) to the DDBJ/EMBL/GenBank databases.  
 Masahira Hattori, The Institute of Physical and Chemical Research  
 (RIKEN), Genomic Sciences Center (GSC), Kitasato Univ., 1-15-1  
 Kitasato, Sagamihara, Kanagawa 228-8555, Japan  
 (E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/  
 Tel:81-42-778-9923, Fax:81-42-778-9924)  
 The sequence is a part of the data (ACCESSION No. AP000165 -  
 AP000173).  
 The sequencing project is supported by Japan Science Technology  
 Corporation (JST) and The Institute of Physical and Chemical  
 Research (RIKEN).  
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 DB 15816 GTTGATGCTGAGAACTTCTACTATGTCAATCCTGTACTCATGGTGTGATGTAATGTTTC 15875  
 QY 181 TCTTTATCATCTGTGGCCATCTCTGTGAGCACTGTGAATTCACAGAGGGGAAACCTCC 240  
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 QY 241 AATGACCCCTACACAGATTCATGAGAGAGTGGCAGAGAAAGTACAGAGCCAAATC 300  
 DB 15936 AATGACCCCTACACAGATTCATGAGAGAGTGGCAGAGAAAGTACAGAGCCAAATC 15995  
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 DB 15996 TTGAATCTAGAAAGATTCGAGAGCCACATCCATGAGAAATTTGGTGGGTTCAAA 16055  
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 DB 16056 ATGTCCCTCTGA 16067  
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 DEFINITION Homo sapiens genomic DNA, chromosome 21q22.1, segment 23/28,  
 complete sequence.  
 ACCESSION AP000052  
 VERSION AP000052.1 GI:3132362  
 KEYWORDS HTG.  
 SOURCE Homo sapiens  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
 REFERENCE 1 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
 TITLE Homo sapiens genomic DNA, chromosome 21q

JOURNAL Published Only In Database (1998) In press  
 REFERENCE 2 (bases 1 to 100000)  
 AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
 TITLE Direct Submission  
 JOURNAL Submitted (11-MAR-1998) to the DDBJ/EMBL/GenBank databases.  
 Masahira Hattori, Kitasato University, Department of Science, JST  
 Sequencing Laboratory; Kitasato 1-15-1, Sagamihara 228, Japan  
 (E-mail:hattori@gsc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/  
 Tel:81-42-778-9923, Fax:81-42-778-9924)  
 This sequence is conducted by Kitasato University JST sequencing  
 Laboratory as a JST sequencing team.  
 Principal Investigator:Yoshiyuki Sakaki Ph.D.  
 Phone: +81-3-5449-5622, Fax: +81-3-5449-5445,  
 sakaki@gsc.riken.go.jp  
 Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D. The  
 sequence is submitted by:human Genome Sequencing In ALIS project of  
 JST  
 Japan Science and Technology Corporation (JST)  
 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0028 Japan  
 For further information about this sequence, including its location  
 and relationship to other sequences, please visit our sequence  
 archive Web site (<http://www.alis.tokyo.jst.go.jp/BSG/top.html>)  
 or send email to [webmaster@www-alis.tokyo.jst.go.jp](mailto:webmaster@www-alis.tokyo.jst.go.jp) ;.  
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 Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
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 DB 80278 ACTTATATGAGCAATTTGGGGCCAGACACAGCTGACAGAGGCGCTCCAGCCAAA 80337  
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 DB 80338 GTTGATGCTGAGAACTTCTACTATGTCAATCCTGTACTCATGGTGTGATGTAATGTTTC 80397  
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 QY 241 AATGACCCCTACACAGATTCATGAGAGAGTGGCAGAGAAAGTACAGAGCCAAATC 300  
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 QY 301 TTGAATCTAGAAAGATTCGAGAGCCACATCCATGAGAAATTTGGTGGGTTCAAA 360  
 DB 80518 TTGAATCTAGAAAGATTCGAGAGCCACATCCATGAGAAATTTGGTGGGTTCAAA 80577  
 QY 361 ATGTCCCTCTGA 372  
 DB 80578 ATGTCCCTCTGA 80589  
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 LOCUS AP000120 100000 bp DNA linear PRI 25-SEP-1999  
 DEFINITION Homo sapiens genomic DNA of 21q22.1, GART and AML related,  
 SLC5A3-f4A4 region, segment 3/8, complete sequence.  
 ACCESSION AP000120  
 VERSION AP000120.1 GI:4730889

KEYWORDS HMG.  
SOURCE Homo sapiens DNA.  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.  
TITLE Homo sapiens 817,199bp genomic DNA of 21q22.1 GART and AML region  
JOURNAL Published Only in Database (1999) In press  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori,M., Yamaguchi,H., Imai,K. and Shimada,J.  
TITLE Direct Submission  
JOURNAL Submitted (15-APR-1999) to the DDBJ/EMBL/Genbank databases. Mita Hirakawa, Japan Science and Technology Corporation (JST), Advanced Databases Department; 5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081, Japan (E-mail:mikasetokyo.jst.go.jp, URL:http://www-alls.tokyo.jst.go.jp/, Tel:01-3-5214-8491, Fax:01-3-5214-8470)  
COMMENT This sequence is conducted by Kitasato University JST sequencing laboratory as a JST sequencing team.  
Principal Investigator:Yoshiyuki Sakaki Ph.D.  
Phone: +81-3-5449-5622, Fax: +81-3-5449-5445, sakaki@hgci.ims.u-tokyo.ac.jp  
Sub-leader: Tadayoshi Shiba Ph.D., Masahira Hattori Ph.D. The sequence is submitted by Human Genome Sequencing in ALIS project of JST.  
Japan Science and Technology Corporation (JST)  
5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081 Japan  
For further information about this sequence, including its location and relationship to other sequences, please visit our sequence archive Web site (http://www-alls.tokyo.jst.go.jp/HGS/) or send email to webmaster@www-alls.tokyo.jst.go.jp.  
Location/Qualifiers  
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LOCUS Homo sapiens genomic DNA, chromosome 21q22.1, D21S226-AML region,  
DEFINITION clone B2344F14-f50E8, segment 3/9, complete sequence.  
ACCESSION AP000167  
VERSION AP000167.1 GI:4827132  
SOURCE Homo sapiens DNA.  
ORGANISM Homo sapiens  
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P., Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Homo sapiens 890,291bp genomic DNA of 21q22.1 (REGION: D21S226-AML CLONE RANGE: B2344F14-f50E8)  
JOURNAL Published Only in Database (1999) In press  
REFERENCE 2 (bases 1 to 100000)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P., Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.  
TITLE Direct Submission  
JOURNAL Submitted (10-MAY-1999) to the DDBJ/EMBL/Genbank databases. Masahira Hattori, The Institute of Physical and Chemical Research (RIKEN), Genomic Sciences Center (GSC), Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555, Japan (E-mail:hattori@gsc.riken.go.jp, URL:http://hgc.riken.go.jp/, Tel:81-42-778-9923, Fax:81-42-778-9924)  
COMMENT E. coli transposon insertion:The present data does not contain E. coli transposon sequences which integrated in the original/previous sequences. We determined the boundary between the insertion and genomic sequences experimentally, removed the insertion sequences, reconstituted the present data. The sequencing project is supported by Japan Science Technology Corporation (JST) and The Institute of Physical and Chemical Research (RIKEN).  
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Best Local Similarity 100.0%; Pred. No. 3.9e-100;  
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

[illegible]

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* URL: http://www.dmb.med.keio.ac.jp/
* GBF, Dept. of Genome Analysis,
* Mascheroder Weg 1, D-38124 Braunschweig, Germany, * e.mail:
* info.genome@gbf.de
* URL: http://genome.gbf.de/
and
* Max-Planck Institute for Molecular Genetics,
* Ihnestrasse 73, D-14195 Berlin, Germany,
* e.mail: info-chr21emo1ogen.mpg.de
* URL: http://chr21.rz-berlin.mpg.de/
AL163264: Submitted (10-Apr-2000).
Location/Qualifiers
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Query Match

100.0%; Score 372; DB 9; Length 340000;

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DB 301487 ACTTATATGACATTTGGCCGCCAGAACACACAGCTGACAGAGAGCCCTCCAGCCAA 301546
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DB 301547 GTTGAATCTGAGAACTTCTACTATGTCATCCTGTACTCTATGATGATGATGTTTC 301606
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DB 301607 TCTTTCATCATGCTGGCCATCCCTGGTGGAGACTGTGAATCCAGAGAGGGGAAACCTGC 301666
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DB 301787 ATGTCCCTCTGA 301798

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RESULT 8
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IMAGE:4481325, mRNA, complete cds.
ACCESSION         BC022699
VERSION           BC022699.1 GI:18490550
KEYWORDS          house mouse.
SOURCE            Mus musculus
ORGANISM          Mus musculus
REFERENCE          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
TITLE             1 (bases 1 to 1664)
JOURNAL            Strausberg/R.

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REMARK  
COMMENT  
NIR-MGC Project URL: <http://mgc.ncl.nih.gov>  
Contact: MGC help desk  
Email: [cgabs-remail.nih.gov](mailto:cgabs-remail.nih.gov)

Tissue Procurement: The Cepho Laboratory  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Sequencing Group at the Stanford Human Genome  
Center, Stanford University School of Medicine, Stanford, CA 94305  
Web site: <http://www-shgc.stanford.edu>  
Contact: (Dickson, Mark) [mdedpaxil.stanford.edu](mailto:mdedpaxil.stanford.edu)  
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,  
R. M.

Clone distribution: MGC clone distribution information can be found  
through the I.M.A.G.E. Consortium/LNL at: <http://image.lnl.gov>  
Series: IRAX Plate: 44 Row: a Column: 22  
This clone was selected for full length sequencing because it  
passed the following selection criteria: Hexamer frequency ORF  
analysis.

FEATURES  
source  
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/organism="Mus musculus"



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/protein_id="AAH2899.1"
/db_xref="GI:1849051"
/translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
LEDSKATIHENLGAFTVSP"

BASE COUNT      502 a      337 c      404 g      421 t
ORIGIN
Query Match      72.0%; Score 268; DB 10; Length 1664;
Best Local Similarity 82.5%; Pred. No. 2.4e-69;
Matches 307; Conservative 0; Mismatches 65; Indels 0; Gaps 0;

QY 1 ATGTCTACTTTATCCAAATTTCACACAGCGCTGGAAGCGTCTCCGAAGATTTTAT 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 90 ATGCCACATTAGCCAAATTGACCCAGACACTGGAGATGCTTCAAAAAGATTTTAT 149
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 ACTTATATGACAAATTGGCGCCAGAACACAAAGCTGACGAAGAGCCCTCCAGCCAA 120
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 150 ACTTATATGACAGCTGGAGAGAGAACACAGCGGAGAGAGAGAGAGAGAGAGAG 209
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 121 GTTGATGCTGAGAACTTCTACTATGTCTACTCTCATGCTGATGATGATGATGAT 180
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    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 210 GTGGATGCGGAGAACTTCTACTAGCTCATCTGCTCATGCTGATGATGATGATG 269
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 181 TCTTTCATCATCTGCGCCATCTCTGTGAGCACTGTGAATCCAAAGAGCGGAACTCC 240
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 270 TCCTTCATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 329
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 241 AATGACCCCTACACAGTACATTTAGAGAGACTGGACAGAAAGTACAAAGAGCCAAATC 300
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 330 CACACACCCCTACACAGTACATTTAGAGAGACTGGACAGAAAGTACAAAGAGCCAAATC 389
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 301 TTGAATCTGAGAAATCGAAGGCCACCATTCATGAGAAATTTGTGCGCTGGGTTCAAA 360
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:1849051"
    /translation="MATLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 390 CTCATCTGGAAGACTCCAGAGGCCACCATTCATGAGAAATTTGTGCGCTGGGTTCAAA 449
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 361 ATGTCCCTCGA 372
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 450 GTGTCAACCTGA 461
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 9
AY050513      372 bp  mRNA  linear  ROD 15-OCT-2001
LOCUS
DEFINITION
Cavia porcellus mink-related peptide 1 mRNA, complete cds.
ACCESSION
AY050513
VERSION
AY050513.1 GI:16151156
KEYWORDS
SOURCE
ORGANISM
domestic guinea pig.
Cavia porcellus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Hystriocognath; Cavidae; Cavia.
REFERENCE
1 (bases 1 to 372)
AUTHORS
Jiang,M., Zhang,M., Liu,J. and Tseng,G.-N.
TITLE
Submitted (12-AUG-2001) Physiology, Virginia Commonwealth
University, 1101 East Marshall Street, Richmond, VA 23298, USA
JOURNAL
Location/Qualifiers
1..372
/organism="Cavia porcellus"
/db_xref="taxon:10141"
/tissue_type="heart"
1..372
CDS
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/note="MIRP1"
/codon_start=1
/product="mink-related peptide 1"
/protein_id="AAH13163.1"
/db_xref="GI:16151157"
/translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
LEDSKATIHENLGAFTVSP"

BASE COUNT      95 a      98 c      105 g      74 t
ORIGIN
Query Match      71.6%; Score 266.4; DB 10; Length 372;
Best Local Similarity 82.3%; Pred. No. 5.2e-69;
Matches 306; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

QY 1 ATGTCTACTTTATCCAAATTTCACACAGCGCTGGAAGCGTCTCCGAAGATTTTAT 60
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    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 1 ATGCCACATTAGCCAAATTGACCCAGACACTGGAGATGCTTCAAAAAGATTTTAT 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 ACTTATATGACAAATTGGCGCCAGAACACAAAGCTGACGAAGAGCCCTCCAGCCAA 120
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    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 61 ACTTATATGACAGCTGGAGAGAGAACACAGCGGAGAGAGAGAGAGAGAGAGAGAG 120
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 121 GTTGATGCTGAGAACTTCTACTATGTCTACTCTCATGCTGATGATGATGATGAT 180
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 121 GTGGATGCGGAGAACTTCTACTAGCTCATCTGCTCATGCTGATGATGATGATGAT 180
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 181 TCTTTCATCATCTGCGCCATCTCTGTGAGCACTGTGAATCCAAAGAGCGGAACTCC 240
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 181 GCCTTCATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 240
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 241 AATGACCCCTACACAGTACATTTAGAGAGACTGGACAGAAAGTACAAAGAGCCAAATC 300
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 241 CAGGACCCCTACACAGTACATTTAGAGAGACTGGACAGAAAGTACAAAGAGCCAAATC 300
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 301 TTGAATCTGAGAAATCGAAGGCCACCATTCATGAGAAATTTGTGCGCTGGGTTCAAA 360
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    /db_xref="GI:16151157"
    /translation="MTTLANLTQLEDAFKFITFYMDSRNRNTTAEQALQARVDAE
    NEYVILVIMWIGMFAYIVAILVSVKSRRESQHPHYQIYVEDMQKYSQILH
    LEDSKATIHENLGAFTVSP"
DB 301 TTGCATCTGGAAGACTCCAGAGGCCACCATTCATGAGAAATTTGTGCGCTGGGTTCAAA 360
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 361 ATGTCCCTCGA 372
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 361 GTGTCAACCTGA 372
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 10
AF071003      468 bp  mRNA  linear  ROD 29-APR-1999
LOCUS
DEFINITION
Rattus norvegicus mink-related peptide 1 mRNA, complete cds.
ACCESSION
AF071003
VERSION
AF071003.1 GI:4704424
KEYWORDS
SOURCE
ORGANISM
Norway rat.
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE
1 (bases 1 to 468)
AUTHORS
Abdolt,G.W., Sestl,F., Splawski,I., Buck,M.E., Lehmann,M.H.,
Timothy,K.W., Keating,M.T. and Goldstein,S.A.
TITLE
MIRP1 forms IKR potassium channels with HERG and is associated with
cardiac arrhythmia
Cell 97 (2), 175-187 (1999)
JOURNAL
99235979
2 (bases 1 to 468)
REFERENCE
Abdolt,G.W., Sestl,F., Buck,M.E. and Goldstein,S.A.N.
AUTHORS
Direct Submission
TITLE
Submitted (05-JUN-1998) Section of Developmental Biology and
Biophysics, Department of Pediatrics and Boyer Center for Molecular
Medicine, Yale University School of Medicine, 295 Congress Avenue,
New Haven, CT 06536, USA
JOURNAL
Location/Qualifiers
1..468
/organism="Rattus norvegicus"
1..468
CDS
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CDs  
/strain="Sprague-Dawley"  
/db\_xref="taxon:10116"  
/sex="male"  
/tissue\_type="heart"  
/disease="10-12 weeks"  
35, 406  
/note="potassium channel subunit; MIRP1"  
/codon\_start=1  
/product="mink-related peptide 1"  
/protein\_id="AD28087.1"  
/db\_xref="GI:4704425"  
/translation="MTTIANLQTLDAKKFVITYWDSRRNTAEQALQARVDAE  
NRYVILYLMVMIGMRAFIYALIVSVSKRREHSODPHQYIVEDMOCKTSQILH  
LEDKATIHENLDAIGFTVSP"  
BASE COUNT 118 a 126 c 131 g 93 t  
ORIGIN

Query Match 71.6%; Score 266.4; DB 10; Length 468;  
Best Local Similarity 82.3%; Pred. No. 5.5e-69;  
Matches 306; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

QY 1 ATGTCTACTTATTCATTTTCACAGACGCTGGAGACGTCCTCCGAGATTTTATT 60  
DB 35 ATGACCACTTATGCACTTATGACGACACCTGGAGATGCTTCAGAAAAGTTTCAT 94  
QY 61 ACTATATGACAAATTTGGCCGACAGACAAACAGCTGAGAGGCCCTCCAGCCAAA 120  
DB 95 ACTATATGACAGCTGAGAGAGAGACAAACAGCCGAGGCTCCAGCCAGA 154  
QY 121 GTTGATGCTGAGACCTTCTATATGTCATCCCTGACCTCATGATGATGTAATGTT 180  
DB 155 GTGATGCGGAGACCTTCTATGTCATCTCTGACCTCATGATGATGCGATGTT 214  
QY 181 TCTTTCATCATCTGGCCATCTGGTGGAGCACTGTGAATTCAGAGAGGGAACACT 240  
DB 215 GCGTCATCTGCTGGCCATCTGTTGAGCACTGTGAATTCAGAGGCGGAGCACT 274  
QY 241 AATGACCCCTACCCAGTACATGTAGAGACCTGGAGAGGAGAAAGTACAGAGCCAA 300  
DB 275 CAGGACCGGTACGACGATGATGAGAGATTTGGAGAGAGGATAGAGCAATC 334  
QY 301 TTGAATCTAGAAATGGAAGGCGACCATTCATGAGAAATTTGGTGGGCTTCAA 360  
DB 335 TTGCATCTGGAAGACTCCAGAGCCACCATCATGAGAACTGGGGGCGAGGGTT 394  
QY 361 ATGTCCCCCTGA 372  
DB 395 GTGTACCCCTGA 406

RESULT 11  
AF329636 215 bp mRNA linear MAN 28-SEP-2001  
LOCUS AF329636  
DEFINITION Oryctolagus cuniculus K+/pacemaker channel beta subunit mirp1  
(kcnk2) mRNA, partial cds.  
ACCESSION AF329636  
VERSION AF329636.1 GI:13194729  
SOURCE rabbit.  
ORGANISM Oryctolagus cuniculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus;  
1 (bases 1 to 215)  
AUTHORS Yu, H., Wu, J., Potapova, I., Wymore, R. T., Holmes, B., Zuckerman, J.,  
Pan, Z., Wang, H., Shi, W., Robinson, R. B., El-Maghrabi, M. R.,  
Benjamin, W., Dixon, J., McKinnon, D., Cohen, I. S. and Wymore, R. S.  
Mink-related peptide 1: A beta subunit for the HCN ion channel  
subunit family enhances expression and speeds activation  
Circ. Res. 88 (12), E84-E87 (2001)  
JOURNAL MEDLINE 21313430  
PUBMED 11420311  
REFERENCE 2 (bases 1 to 215)  
AUTHORS Wymore, R. T., Holmes, B. A., Wymore, R. S., Yu, H., Wu, J., Potapova, I.,

Zuckerman, J., Pan, Z., Wang, H., Shi, W., Robinson, R., El-Maghrabi, R.,  
Benjamin, W., Dixon, J. E., McKinnon, D. and Cohen, I. S.  
Direct Submission  
Submitted (15-DEC-2000) Biology, The University of Tulsa, 600 S.  
College Av., Tulsa, OK 74104-3169, USA  
Location/Qualifiers  
FEATURES  
source  
1..215  
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/db\_xref="taxon:9986"  
<1..>215  
/gene="kcnk2"  
<1..>215  
/gene="kcnk2"  
/codon\_start=3  
/product="K+/pacemaker channel beta subunit mirp1"  
/protein\_id="AK15527.1"  
/db\_xref="GI:13194730"  
/translation="AENRYVILYLMVMIGMRAFIYALIVSVSKRREHSNDYHQ  
YIVEDMOCKTSQILHFEAKRTIEN"  
BASE COUNT 60 a 58 c 51 g 46 t  
ORIGIN

Query Match 50.1%; Score 186.2; DB 4; Length 215;  
Best Local Similarity 91.6%; Pred. No. 5.1e-45;  
Matches 197; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

QY 125 ATGCTGAACTTCTACTATGATCATCTGATCTACCTCATGATGATGTAATGTTCT 184  
DB 1 ATGCGGAACTTCTACTATGATCATCTGATCTACCTCATGATGATGATGATGTTCT 60  
QY 185 TCATCATCTGCGCATCTGCTGAGAGCACTGTGAATTCAGAGAGCCGGAACCTCCAA 244  
DB 61 TCATCATCTGCGCATCTGCTGAGAGCACTGTGAATTCAGAGAGCCGGAACCTCCAA 120  
QY 245 ACCCTACACCACTGATGATGAGAGACTGCGAGAGAAAGTACAGAGCCAAATCTGA 304  
DB 121 ACCCTACACCACTGATGATGAGAGACTGCGAGAGAAAGTACAGAGCCAAATCTGA 180  
QY 305 ATCTAGAGAAATGGAAGGCGACCATTCATGAGAAC 339  
DB 181 ATTTGGAAGAGCAAGGCGACCATTCATGAGAAC 215

RESULT 12  
AF387764 228 bp mRNA linear MAN 08-NOV-2001  
LOCUS AF387764  
DEFINITION Equus caballus mink related peptide 1-like mRNA, partial sequence.  
ACCESSION AF387764  
VERSION AF387764.1 GI:16797919  
KEYWORDS  
SOURCE horse.  
ORGANISM Equus caballus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Perissodactyla; Equidae; Equus;  
1 (bases 1 to 228)  
AUTHORS Finley, M. R., Mitchell, K. E., Mitchell, B. M. and Freeman, L. C.  
Post-translational modification of ERG in horse heart  
FASEB J. 15, A1135 (2001)  
REFERENCE 2 (bases 1 to 228)  
AUTHORS Finley, M. R., Li, Y. and Freeman, L. C.  
JOURNAL Direct Submission  
Submitted (31-MAY-2001) Department of Anatomy and Physiology,  
Kansas State University, 1600 Denison Ave, Manhattan, KS 66506, USA  
NCBI staff are still waiting for submitters to provide appropriate  
coding sequence information.  
FEATURES  
source  
1..228  
/organism="Equus caballus"  
/db\_xref="taxon:9796"  
/tissue\_type="atrium"  
44..>228  
/note="similar to mink related peptide 1; potassium  
channel accessory protein; MIRP1, KCN2"

BASE COUNT	68 a	57 c	51 g	51 t	1 others
ORIGIN					
Query Match	47.9%	Score 178.2;	DB 4;	Length 228;	
Best Local Similarity	89.0%;	Pred. No. 1.3e-42;			
Matches 203; Conservative	0;	Mismatches 24;	Indels 1;	Gaps 1;	
Oy	124	GATGCTAGAACTTGTACTATGTCATCCTGTACCTCATCGT-GATGATTGGAAATGTTCTC	182		
Dd	1	GATGGGGAACCTTCTACTACGTCTCTTGACCATTAGCATGATGTAATGGAAATGTTCTC	60		
Oy	183	TTCATCATCGGGCCATCTCGTGAGAGCACTGTGAATCCAAAGAGACGGGAACACTCCAA	242		
Dd	61	TTTTCTCATTTGAGGCATCTGTTGAGAGAGGAGTGAATCCAAAGAGACGAACAACCTCAA	120		
Oy	243	TGACCCCTACCAACAGTACATTGTAGAGAGTGGGAGAGAAAATGACCAAGACCTT	302		
Dd	121	CGACCCCTACCAACAGTACATCTGTAAGAGACTGGCAAGAAATACAGAGAATCAATTTT	180		
Oy	303	GAATCTAGAGAANTCGAAGGCCACCATCATGAGAACATTTGGTGGGC	350		
Dd	181	CAATCTAGAGAACCAAGGCCACCATCATGANAACCTTGGCGGGC	228		
RESULT 13					
RABIPCS	534 bp	DNA	linear	MAM 16-Oct-2001	
LOCUS	Oryctolagus cuniculus cardiac delayed rectifier potassium channel				
DEFINITION	protein gene, complete cds.				
ACCESSION	L41659				
VERSION	L41659.1	GI:1246305			
KEYWORDS					
SOURCE	rabbit.				
ORGANISM	Oryctolagus cuniculus				
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;					
Mammalia; Euthera; Lagomorpha; Leporidae; Oryctolagus.					
REFERENCE	1 (bases 1 to 534)				
AUTHORS	Salate,J.J., Jurkiewicz,N.K., Jow,B., Polander,K., Guinasso,P.J. Jr., Raynor,B., Swanson,R. and Fermil,B.				
TITLE	IK of rabbit ventricle is composed of two currents: evidence for IKs				
JOURNAL	Am. J. Physiol. 271 (6), H2477-H2489 (1996)				
MEDLINE	9751310				
PUBMED	8997308				
FEATURES	Location/Qualifiers				
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	/strain="New Zealand White"				
	/db_xref="taxon:9986"				
	/clone="A1"				
	/clone_1id="Clontech genomic DNA library in EMBL3 SP6/T7"				
	/dev_stage="adult"				
	/note="breed: New Zealand White"				
CDS	133..525				
	/function="cardiac potassium channel"				
	/note="Isk protein"				
	/codon_start=1				
	/product="cardiac delayed rectifier potassium channel protein"				
	/protein_id="AA93505.1"				
	/db_xref="GI:1246306"				
	/translation="MPPNATAVPPLTLTGEEYAHQGSSEATSLARGLPDGDKME ALVIAIVGFGEFTGLIMLSYIRSKQLKLESHDPFNYIEANDQERDRAVFQRVLE SCRRCYVENLQNAVHPDTHLPKLPS"				
BASE COUNT	105 a	175 c	149 g	105 t	
ORIGIN					
Query Match	15.1%;	Score 56;	DB 4;	Length 534;	
Best Local Similarity	66.7%;	Pred. No. 6.6e-06;			
Matches 96; Conservative	0;	Mismatches 45;	Indels 3;	Gaps 1;	
Oy	151	CTGTACCTCAGCGTAGATTTGGAAATGTTCTCTTTCATCATCATCGGCGCATCCTGTGTAGC	210		

QY	211	ACTGTGAATCCCAAGAGACGGGAGACATCCATGACCCCTACACCACTAGATTGTAG--	268
Db	328	TACATCCGCTCCACAGAACTGGAGACATCGCATGACCCCTTCAACGTGTACATTGAGCC	387
QY	269	-AGGACTGGCAGGAAAAGTACAG 291	
Db	388	AACGACTGGCAGGAAAAGACAG 411	
RESULT 14			
LOCUS	GP1ISK	750 bp	linear
DEFINITION	Cavia cobaya potassium channel mRNA, complete cds.		ROD 11-MAY-1994
ACCESSION	L20462		
VERSION	L20462.1		
KEYWORDS	potassium channel.		
SOURCE	Cavia cobaya adult cardiac muscle cDNA to mRNA.		
ORGANISM	Cavia porcellus		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Hystriognathli; Caviidae; Cavia.		
AUTHORS	1 (bases 1 to 750)		
TITLE	Zhang,J., Jurkiewicz,N.K., Folander,K., Lazarides,E., Salata,J.J. and Swanson,R.		
JOURNAL	K+ currents expressed from the guinea pig cardiac IsK protein are enhanced by activators of protein kinase C		
MEDLINE	Proc. Natl. Acad. Sci. U.S.A. 91, 1766-1770 (1994)		
FEATURES	94173910		
source	Location/Qualifiers		
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	/organism="Cavia porcellus"		
	/db_xref="taxon:10141"		
	/cell_type="myocyte"		
	/tissue_type="cardiac muscle"		
	/dev_stage="adult"		
	63..440		
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	/protein_id="AA02394.1"		
	/db_xref="GI:484141"		
	/translation="MLPNSVAVMPELTTWGTVOYPSNNSAGLARSPLRDOCKLEA		
	LYLIMLGFRRFFTLGIMLSYRKKLEHSDPNVYIESDPMENKAPFQARVLENN		
	CRSCCVIENQLTVEQPNITLPEL		
CDS			
BASE COUNT	189 a 199 c 174 g 188 t		
ORIGIN			
Query Match	14.8%; Score 55; DB 10; Length 750;		
Best Local Similarity	66.4%; Pred. No. 1.4e-05;		
Matches 95; Conservative	0; Mismatches 45; Indels 3; Gaps 1;		
QY	151	CTGTACCTCATGGTATGTAATGTCTCTTCATCATCATGTCGGCCATCCTGGTGAGC 210	
Db	195	CTCTACATCTCTATGTGTGTGGCTTCTTTGTTCTTCTACCTGTGGCATCTGCTGAGC 254	
QY	211	ACTGTGAATCCCAAGAGACGGGAGACATCCATGACCCCTACACCACTAGATTG--TA 267	
Db	255	TATATTCATCCCAAGAACTGGAGACATCGACGACCCGTTCAACGTGTACATCGAGTCA 314	
QY	268	GAGGACTGGCAGGAAAAGTACAA 290	
Db	315	GACACCTGGCAGGAAAAGACAA 337	
RESULT 15			
LOCUS	AF135188	390 bp	mRNA
DEFINITION	Human sapiens delayed rectifier potassium channel subunit IsK mRNA, complete cds.		linear
ACCESSION	AF135188		
VERSION	AF135188.1		
KEYWORDS	GI:4583498		
SOURCE	human.		
ORGANISM	Homo sapiens		





XX WIPI: 2001-488901/53.  
 DR Human genome-derived single exon nucleic acid probes useful for  
 XX analyzing gene expression in human cervical epithelial cells -  
 PT  
 XX  
 PS Claim 25; SEQ ID No 14365; 487pp; English.  
 XX  
 CC The present invention relates to human single exon nucleic acid probes  
 CC (SENPs). The present sequence is one such probe. The SENPs are derived  
 CC from human HeLa cells. The SENPs can be used to produce a single exon  
 CC microarray, which can be used for measuring human gene expression in a  
 CC sample derived from human cervical epithelial cells. By measuring gene  
 CC expression, the probes are therefore useful in grading and/or staging  
 CC of diseases of the cervix, notably cervical cancer.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pcl\_sequences.  
 XX  
 SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;  
 Query Match 100.0%; Score 372; DB 22; Length 372;  
 Best Local Similarity 100.0%; Pred. No. 1.6e-102;  
 Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ATGCTACTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCGAGATTTTATT 60  
 DB 1 ATGCTACTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCGAGATTTTATT 60  
 QY 61 ACTATATGACCAATTTGGCCGACAGACACACAGCTGAGCAAGGCGCTCCAGCCAAA 120  
 DB 61 ACTATATGACCAATTTGGCCGACAGACACACAGCTGAGCAAGGCGCTCCAGCCAAA 120  
 QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGATGATTTGAAATGTTTC 180  
 DB 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGATGATTTGAAATGTTTC 180  
 QY 181 TCTTTCATCATCCGTGGCCATCTCTGTGACACTGTGAATTCAGAGAGCGGAACACTCC 240  
 DB 181 TCTTTCATCATCCGTGGCCATCTCTGTGACACTGTGAATTCAGAGAGCGGAACACTCC 240  
 QY 241 AATGACCCCTACACAGTACATTTGTAGAGACTGAGCAAGGAAAAGTACAAAGAGCCAAATC 300  
 DB 241 AATGACCCCTACACAGTACATTTGTAGAGACTGAGCAAGGAAAAGTACAAAGAGCCAAATC 300  
 QY 301 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAACATTTGGTGGCTGGTTCAAA 360  
 DB 301 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAACATTTGGTGGCTGGTTCAAA 360  
 QY 361 ATGTCCCTCTGA 372  
 DB 361 ATGTCCCTCTGA 372  
 RESULT 2  
 AA109965  
 ID AA109965 standard; DNA; 372 BP.  
 XX  
 AC AA109965;  
 XX  
 DT 09-OCT-2001 (first entry)  
 XX  
 DE Probe #9956 used to measure gene expression in human breast sample.  
 XX  
 KW Probe: human; breast disease; breast cancer; development disorder; ss;  
 KW inflammatory disease; proliferative breast disease; non-carcinoma tumour.  
 XX  
 OS Homo sapiens.  
 XX  
 PN W0200157270-A2.  
 XX  
 PD 09-AUG-2001.  
 XX

PF 29-JAN-2001; 2001WO-US00661.  
 XX  
 PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632266.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024263.  
 XX  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 PI Penn SG, Hanzel DK, Chen W, Rank DR;  
 XX  
 XX WIPI: 2001-476286/51.  
 XX  
 PT Novel single exon nucleic acid probe used to measuring gene expression  
 PT in a human breast -  
 CC  
 PS Claim 25; SEQ ID No 9956; 322pp; English.  
 XX  
 CC The present invention relates to novel single exon nucleic acid probes.  
 CC The present sequence is one such probe. The probes are useful for  
 CC measuring human gene expression in a human breast sample, where the probe  
 CC hybridises at high stringency to a nucleic acid expressed in the human  
 CC breast. The probes are useful for predicting, diagnosing, grading,  
 CC staging, monitoring and prognosing diseases of the human breast,  
 CC particularly those diseases with polygenic aetiology. The diseases  
 CC include: breast cancer, disorders of development, inflammatory diseases  
 CC of the breast, fibrocystic changes, proliferative breast disease and  
 CC non-carcinoma tumours.  
 CC Note: The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pcl\_sequences.  
 XX  
 SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;  
 Query Match 100.0%; Score 372; DB 22; Length 372;  
 Best Local Similarity 100.0%; Pred. No. 1.6e-102;  
 Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ATGCTACTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCGAGATTTTATT 60  
 DB 1 ATGCTACTTATTCATTTTCACACAGACGCTGGAAGACGCTTCCGAGATTTTATT 60  
 QY 61 ACTATATGACCAATTTGGCCGACAGACACACAGCTGAGCAAGGCGCTCCAGCCAAA 120  
 DB 61 ACTATATGACCAATTTGGCCGACAGACACACAGCTGAGCAAGGCGCTCCAGCCAAA 120  
 QY 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGATGATTTGAAATGTTTC 180  
 DB 121 GTTGATGCTGAGAACTTCTACTATGTCATCCTGTACCTCATGTGATGATTTGAAATGTTTC 180  
 QY 181 TCTTTCATCATCCGTGGCCATCTCTGTGACACTGTGAATTCAGAGAGCGGAACACTCC 240  
 DB 181 TCTTTCATCATCCGTGGCCATCTCTGTGACACTGTGAATTCAGAGAGCGGAACACTCC 240  
 QY 241 AATGACCCCTACACAGTACATTTGTAGAGACTGAGCAAGGAAAAGTACAAAGAGCCAAATC 300  
 DB 241 AATGACCCCTACACAGTACATTTGTAGAGACTGAGCAAGGAAAAGTACAAAGAGCCAAATC 300  
 QY 301 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAACATTTGGTGGCTGGTTCAAA 360  
 DB 301 TTGAATCTAGAAAGATCGAAGGCCACATCCATGAGAAACATTTGGTGGCTGGTTCAAA 360  
 QY 361 ATGTCCCTCTGA 372  
 DB 361 ATGTCCCTCTGA 372  
 RESULT 3  
 AAS00245  
 ID AAS00245 standard; DNA; 372 BP.

```

XX AC AAS0245;
XX 10-MAR-2001 (first entry)
XX DE Human potassium channel regulatory protein, Mink2, DNA sequence.
XX KM Human; Mink2; potassium channel; cardiac arrhythmia; hypertension; ds;
XX KM angina; asthma; diabetes; renal insufficiency; urinary incontinence;
XX KM irritable colon; epilepsy; cerebrovascular ischemia; autoimmune disease.
XX OS Homo sapiens.
XX FH Key 1.372
XX FT CDS /tag= a
XX FT /product= "MINK2 potassium channel protein"
XX PN WO200114403-A1.
XX PD 01-MAR-2001.
XX PF 18-AUG-2000; 2000WO-US22799.
XX PR 20-AUG-1999; 99US-0379201.
XX PA (UYCA-) UNIV CASE WESTERN RESERVE.
XX PI Flicker E, Wible B, Brown AM;
XX PI WPI; 2001-218424/22.
XX DR P-PSDB; AAN00215.
XX PT Novel potassium channel gene termed Mink2 encoding potassium channel
XX PT regulatory protein, useful for screening compounds that are useful for
XX PT treating diseases caused by aberrant potassium activity -
XX PS Claim 1; Fig 9; 39pp; English.
XX CC The sequence represents the coding sequence of human potassium channel
XX CC regulatory protein, Mink2. Mink2 sequence is useful for producing a
XX CC potassium channel regulatory protein useful for in vitro or in vivo
XX CC screening of agonistic or antagonistic compounds that are useful for
XX CC treating diseases caused by aberrant potassium activity, such as human
XX CC cardiac arrhythmias, hypertension, angina, asthma, diabetes, renal
XX CC insufficiency, urinary incontinence, irritable colon, epilepsy,
XX CC cerebrovascular ischemia, and autoimmune disease.
XX SQ Sequence 372 BP; 110 A; 90 C; 82 G; 90 T; 0 other;

Query Match 100.0%; Score 372; DB 22; Length 372;
Best Local Similarity 100.0%; Pred. No. 1.6e-102;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGTTACTTATTCATTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 60
DB 1 ATGTTACTTATTCATTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 60
QY 61 ACTTATATGACAAATTTGGCCGACAAACACACAGCTGACCAAGAGCCCTCCAA 120
DB 61 ACTTATATGACAAATTTGGCCGACAAACACACAGCTGACCAAGAGCCCTCCAA 120
QY 121 GTTGATGCTGAGAACTTCTACATGTCATCCGTCATCCATGATGATTTGAT 180
DB 121 GTTGATGCTGAGAACTTCTACATGTCATCCGTCATCCATGATGATTTGAT 180
QY 181 TCTTTCATCATCGTGGCCATCCTGCTGAGCACTGTGAATCCAAAGAGGGAACACT 240
DB 181 TCTTTCATCATCGTGGCCATCCTGCTGAGCACTGTGAATCCAAAGAGGGAACACT 240
QY 241 AATGACCCCTACACACAGTACATTGTAGAGAGCTGCGACGAAAGATACAAAGCA 300
DB 241 AATGACCCCTACACACAGTACATTGTAGAGAGCTGCGACGAAAGATACAAAGCA 300

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QY 301 TTGAATCTAGAAGATCGAAGCCACCATCCATGAGAACATTTGGTGGCTGGTCA 360
DB 301 TTGAATCTAGAAGATCGAAGCCACCATCCATGAGAACATTTGGTGGCTGGTCA 360
QY 361 ATGTCCCCCTGA 372
DB 361 ATGTCCCCCTGA 372

RESULT 4
ID AAF80269 standard; DNA; 471 BP.
XX AAF80269;
AC AAF80269;
XX 29-JUN-2001 (first entry)
XX DE Nucleotide sequence of human potassium channel subunit Isk2.
XX KM Human; potassium channel; Isk2; gene therapy; gastric motility;
XX KM gastric acid secretion; anti-arrhythmic agent; myocardial infarction; ss.
XX OS Homo sapiens.
XX FH Key Location/Qualifiers
XX FT CDS 79..450
XX FT /*tag= a
XX FT /product= "potassium channel subunit Isk2"
XX PN WO200127246-A1.
XX PD 19-APR-2001.
XX PF 10-OCT-2000; 2000WO-US28014.
XX PR 12-OCT-1999; 99US-0158781.
XX PA (MERI ) MERCK & CO INC.
XX PI Swanson RJ, Liu Y, Folander K;
XX PI WPI; 2001-273764/28.
XX DR P-PSDB; AAB67800.
XX PT New DNA encoding the Isk2 potassium channel subunit, useful e.g. for
XX PT detecting mutations and screening for therapeutic agents -
XX PS Claim 3; Fig 1A; 46pp; English.
XX CC The present sequence encodes a human potassium channel subunit,
XX CC designated Isk2. The Isk2 polynucleotide, and derived probes, are
XX CC used diagnostically to detect mutations in the Isk2 gene, to determine
XX CC levels of mRNA expression and to isolate homologous sequences; for
XX CC recombinant expression of Isk2; in gene therapy to increase potassium
XX CC channel activity and to generate transgenic animals, as models and
XX CC for drug screening. Recombinant Isk2 is used for studying biochemical
XX CC activity of Isk2 and its role in disorders of gastric motility and
XX CC gastric acid secretion, and to raise specific antibodies. Isk2
XX CC modulators are potentially useful for treating diseases associated with
XX CC increased or reduced potassium channel activity, e.g. as
XX CC anti-arrhythmic agents for treating myocardial infarction and as
XX CC regulators of gastric acid secretion.
XX SQ Sequence 471 BP; 143 A; 110 C; 103 G; 115 T; 0 other;

Query Match 100.0%; Score 372; DB 22; Length 471;
Best Local Similarity 100.0%; Pred. No. 1.8e-102;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGTTACTTATTCATTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 60
DB 79 ATGTTACTTATTCATTCACACAGACGCTGGAAGCGTTCGGAAGATTTTAT 138

```

QY 61 ACTATATGACAAATTGGCGCCAGAACACACAGCTGAGCAAGGCGCCCTCCAGCCAAA 120  
DB 139 ACTTATATGACAAATTGGCGCCAGAACACACAGCTGAGCAAGGCGCCCTCCAGCCAAA 198  
QY 121 GTTGATGCTGAGAACTTCTACTATGTCTCTGTACCTGCATGATGATGATGATGATGTC 180  
DB 199 GTTGATGCTGAGAACTTCTACTATGTCTCTGTACCTGCATGATGATGATGATGATGTC 258  
QY 181 TCTTTCATCATGTCGGCCATCTGCTGTGAGCACTGTGAAATCCAGAGAGCGGAACACTCC 240  
DB 259 TCTTTCATCATGTCGGCCATCTGCTGTGAGCACTGTGAAATCCAGAGAGCGGAACACTCC 318  
QY 241 AATGACCCCTACACCACTACATGTTAGAGAGCTGGCAGAGAAAGTACAGAGCCAAATC 300  
DB 319 AATGACCCCTACACCACTACATGTTAGAGAGCTGGCAGAGAAAGTACAGAGCCAAATC 378  
QY 301 TTGAATCTAGAAAGATGCAAGGCGCCATCATGATGAAATGTCGGCGCTGCTCAAA 360  
DB 379 TTGAATCTAGAAAGATGCAAGGCGCCATCATGATGAAATGTCGGCGCTGCTCAAA 438  
QY 361 ATGTCCCCCTGA 372  
DB 439 ATGTCCCCCTGA 450

RESULT 5  
ABA09192 standard; cDNA: 600 BP.  
ID ABA09192  
XX ABA09192:  
AC 11-JAN-2002 (first entry)  
DT Human M1RP1 homologue-encoding cDNA, SEQ ID NO:968.  
XX  
DE Human M1RP1 homologue-encoding cDNA, SEQ ID NO:968.  
XX  
XX Human: cytokine; cell proliferation; cell differentiation; growth factor;  
KW hemotopolesis regulation; tissue growth; immunomodulator; activin;  
KW inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;  
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;  
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;  
KW chronic inflammatory condition; proliferative retinopathy;  
KW atherosclerosis; coronary heart disease; arterial ischaemia;  
KW bone disorder; osteoporosis; vascular growth disorder;  
KW tissue regeneration; wound healing; infection; immune disorder;  
KW cell culture; drug screening; gene therapy; antiinflammatory;  
KW antiaesthetic; antiallergic; haemostatic; antiarteriosclerotic;  
KW cytosolic; osteopathic; vasotropic; cardiac; virucide; antibacterial;  
KW antifungal; vulnery; antitumor; ss.  
XX  
OS Homo sapiens.  
XX  
XX WO200157188-A2.  
XX  
XX 09-AUG-2001.  
XX  
XX 05-FEB-2001; 2001MO-US03800.  
XX  
XX 03-FEB-2000; 2000US-0496914.  
XX 27-APR-2000; 2000US-0560875.  
XX  
XX (HYSE-) HYSEQ INC.  
XX  
XX Tang YT, Liu C, Drmanac RT;  
XX  
XX WPI: 2001-457740/49.  
XX  
XX DR P-PSDB; ABB11948.  
XX  
XX Human proteins and DNA encoding sequences useful for preventing,  
XX treating or ameliorating a medical condition in a mammalian subject  
XX e.g. arthritis and cancer -  
XX  
PS Claim 1; Page 826; 1963pp; English.

XX  
CC Sequences ABB10981-ABB12330 r'present 1350 novel human polypeptides, and  
CC sequences ABA08225-ABA09574 r'present nucleic acids encoding them. The  
CC invention also relates to vectors and recombinant host cells comprising a  
CC nucleotide of the invention, methods of producing the novel polypeptides,  
CC antibodies against the polypeptides, methods of detecting the nucleotides  
CC or polypeptides in a sample, and methods of identifying compounds which  
CC bind to polypeptides of the invention. Although novel, many of the  
CC polypeptides of the invention have homology to known proteins, thereby  
CC giving an insight into their probable biological activities, and hence  
CC potential therapeutic applications. The polypeptides of the invention may  
CC have various activities, including cytokine, cell proliferation or cell  
CC differentiation activities; stem cell growth factor activity;  
CC haematopoiesis regulatory activity; tissue growth activity;  
CC immunomodulatory activity; activin- or inhibin-related activities;  
CC chemotactic or chemokinetic activities; haemostatic, thrombotic or  
CC thrombolytic activities; receptor or ligand activities; or may be  
CC involved in oncogenesis, cancer cell proliferation or metastasis.  
CC Depending on their biological activities, polypeptides and nucleotides of  
CC the invention are useful for preventing, treating or ameliorating medical  
CC conditions, e.g., by protein or gene therapy. Such conditions include  
CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell  
CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),  
CC proliferative retinopathy, atherosclerosis, coronary heart disease,  
CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal  
CC vascular growth. Polypeptides involved with tissue regeneration and  
CC repair (or nucleic acids encoding them) may be used to promote wound  
CC healing (e.g., of burns, incisions and ulcers), while those with  
CC immunomodulatory activities may be used in the treatment of viral,  
CC bacterial and fungal infections in addition to immune disorders.  
CC Polypeptides with growth factor activity may be used in cell cultures to  
CC promote cell growth. For example, such polypeptides may be used to  
CC manipulate stem cells in culture to give rise to neuroepithelial cells  
CC that can be used to augment or replace cells damaged by illness,  
CC autoimmune disease or accidental damage. The polypeptides and nucleotides  
CC may also be used in the diagnosis of the above conditions, and in drug  
CC screening techniques. The present sequence represents a cDNA encoding a  
CC novel human polypeptide of the invention.  
XX  
SQ Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;  
XX  
XX  
XX Query Match 100.0%; Score 372; DB 22; Length 600;  
XX Best Local Similarity 100.0%; Pred. No. 2e-102;  
XX Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
QY 1 ATGTCTACTTATTCATTTTACACAGAGCGCTGGAAGCGTCTCCGAAGATTTTATT 60  
DB 38 ATGTCTACTTATTCATTTTACACAGAGCGCTGGAAGCGTCTCCGAAGATTTTATT 97  
QY 61 ACTTATATGACAAATTGGCGCCAGAACACACAGCTGAGCAAGGCGCCCTCCAGCCAAA 120  
DB 98 ACTTATATGACAAATTGGCGCCAGAACACACAGCTGAGCAAGGCGCCCTCCAGCCAAA 157  
QY 121 GTTGATGCTGAGAACTTCTACTATGTCTCTGTACCTGCATGATGATGATGATGATGTC 180  
DB 158 GTTGATGCTGAGAACTTCTACTATGTCTCTGTACCTGCATGATGATGATGATGATGTC 217  
QY 181 TCTTTCATCATGTCGGCCATCTGCTGTGAGCACTGTGAAATCCAGAGAGCGGAACACTCC 240  
DB 218 TCTTTCATCATGTCGGCCATCTGCTGTGAGCACTGTGAAATCCAGAGAGCGGAACACTCC 277  
QY 241 AATGACCCCTACACCACTACATGTTAGAGAGCTGGCAGAGAAAGTACAGAGCCAAATC 300  
DB 278 AATGACCCCTACACCACTACATGTTAGAGAGCTGGCAGAGAAAGTACAGAGCCAAATC 337  
QY 301 TTGAATCTAGAAAGATGCAAGGCGCCATCATGATGAAATGTCGGCGCTGCTCAAA 360  
DB 338 TTGAATCTAGAAAGATGCAAGGCGCCATCATGATGAAATGTCGGCGCTGCTCAAA 397  
QY 361 ATGTCCCCCTGA 372  
DB 398 ATGTCCCCCTGA 409



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RESULT 6
AAK52645
ID AAK52645 standard; cDNA; 600 BP.
XX
XX AAK52645;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 2174.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation; ss.
XX
OS Homo sapiens.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US04098.
XX
PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
XX (HYSE-) HYSEQ INC.
XX
PI Tang YF, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QJ, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX
XX WPI; 2001-476283/51.
XX
XX P-PSDB; AAM79512.
XX
PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
XX
XX Claim 1; Page 4539-4540; 6221pp; English.
XX
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
CC production of other cytokines in other cell populations. The
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
CC peptide therapy. The polypeptides have various cytokine-like activities,
CC e.g. stem cell growth factor activity, haematopoiesis regulating
CC activity, tissue growth factor activity, immunomodulatory activity and
CC activin/inhibin activity and may be useful in the diagnosis and/or
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
CC inflammation.
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666
CC (AAM80020) are omitted as the relevant pages from the sequence listing
CC were missing at the time of publication.
XX
SO Sequence 600 BP; 187 A; 133 C; 130 G; 144 T; 6 other;
Query Match 100.0%; Score 372; DB 22; Length 600;
Best Local Similarity 100.0%; Pred. No. 2e-102;
Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGTCTACTTATCCAAATTTACACAGAGCGCTGGAAGAGCTCTCCGAAGGATTTTATT 60
DB 38 ATGTCTACTTATCCAAATTTACACAGAGCGCTGGAAGAGCTCTCCGAAGGATTTTATT 97
QY 61 ACTTATATGAGCAATTGGCGCCAGAACACAAAGCTGAGCAAGAGGCGCTCCAAAGCCAAA 120
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DB 98 ACTTATATGAGCAATTGGCGCCAGAACACAAAGCTGAGCAAGAGGCGCTCCAAAGCCAAA 157
QY 121 GTTGATGCTGAGACTTCTACTATGTATCTCTGTACTCTCATGTGATGAAATGTC 180
DB 158 GTTGATGCTGAGACTTCTACTATGTATCTCTGTACTCTCATGTGATGAAATGTC 217
QY 181 TCTTTCATCATGCTGGCCATCTGCTGAGCACTGTGAATCCAAAGAGCGGAACATGCC 240
DB 218 TCTTTCATCATGCTGGCCATCTGCTGAGCACTGTGAATCCAAAGAGCGGAACATGCC 277
QY 241 AATGACCCCTACACACAGTATCTGTAGAGCACTGGCAGAAAGTCAACAGCCAAATC 300
DB 278 AATGACCCCTACACACAGTATCTGTAGAGCACTGGCAGAAAGTCAACAGCCAAATC 337
QY 301 TTGAATCTAGAGAAATCGAAGGCCACCATCATGAGAAATTTGTGCGGTGTTCAA 360
DB 338 TTGAATCTAGAGAAATCGAAGGCCACCATCATGAGAAATTTGTGCGGTGTTCAA 397
QY 361 ATGTCCCTCTGA 372
DB 398 ATGTCCCTCTGA 409
RESULT 7
AAK51661
ID AAK51661 standard; cDNA; 655 BP.
XX
XX AAK51661;
XX
AC AAK51661;
XX
DT 06-NOV-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 206.
XX
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW tissue growth factor; immunomodulatory; cancer; leukaemia;
KW nervous system disorder; arthritis; inflammation; ss.
XX
XX Homo sapiens.
XX
XX WO200157190-A2.
XX
PN WO200157190-A2.
XX
PD 09-AUG-2001.
XX
PF 05-FEB-2001; 2001WO-US04098.
XX
PR 03-FEB-2000; 2000US-0496914.
PR 27-APR-2000; 2000US-0560875.
PR 20-JUN-2000; 2000US-0598075.
PR 19-JUL-2000; 2000US-0620325.
PR 01-SEP-2000; 2000US-0654936.
PR 15-SEP-2000; 2000US-0663561.
PR 20-OCT-2000; 2000US-0693325.
PR 30-NOV-2000; 2000US-0728422.
XX
XX (HYSE-) HYSEQ INC.
XX
PI Tang YF, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI Zhao QJ, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;
XX
XX WPI; 2001-476283/51.
XX
XX P-PSDB; AAM78528.
XX
PT Nucleic acids encoding polypeptides with cytokine-like activities,
PT useful in diagnosis and gene therapy -
XX
XX Claim 1; Page 1024; 6221pp; English.
XX
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to
CC cytokine, cell proliferation or cell differentiation or which may induce
```

CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAB60020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX

SO Sequence 655 BP; 196 A; 154 C; 146 G; 153 T; 6 other;

Query Match 100.0%; Score 372; DB 22; Length 655;  
Best Local Similarity 100.0%; Pred. No. 2.1e-102;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGTCTACTTATTCATTTTCACACAGACGCTGGAGACGCTCCGAGAGATTATTAT 60  
DB 93 ATGTCTACTTATTCATTTTCACACAGACGCTGGAGACGCTCCGAGAGATTATTAT 152  
OY 61 ACTTATATGACAAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 120  
DB 153 ACTTATATGACAAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 212  
OY 121 GTTATATGCTGAGAACTTCTACTATATCTCTGACTCATGTGATGATTTGAAATGTC 180  
DB 213 GTTATATGCTGAGAACTTCTACTATATCTCTGACTCATGTGATGATTTGAAATGTC 272  
OY 181 TCTTTCATCATCGTGGCCATCTGTGAGACAGCTGAAATCCAAAGACGGAACTCC 240  
DB 273 TCTTTCATCATCGTGGCCATCTGTGAGACAGCTGAAATCCAAAGACGGAACTCC 332  
OY 241 AATGACCCCTACACACAGTACATTTAGAGAGACTGGCAGAGAAAAGTCAAGAGCCAAATC 300  
DB 333 AATGACCCCTACACACAGTACATTTAGAGAGACTGGCAGAGAAAAGTCAAGAGCCAAATC 392  
OY 301 TTGAAATCTAGAAATGGAAGCCACCATTCATGAGAAACATTTGGCGCTGGTTCAAA 360  
DB 393 TTGAAATCTAGAAATGGAAGCCACCATTCATGAGAAACATTTGGCGCTGGTTCAAA 452  
OY 361 ATGTCCCTCTGA 372  
DB 453 ATGTCCCTCTGA 464

RESULT 8  
AAC64071  
ID AAC64071 standard; cDNA; 732 BP.

XX AAC64071;

DT 19-FEB-2001 (first entry)

DE Human potassium channel protein KCNE2 (MiRP1) cDNA, SEQ ID NO:1.

XX Human; KCNE2; MiRP1; potassium channel protein; KCNE1-related;  
XX MiRP-related; long QT syndrome; cardiac arrhythmia;  
XX drug screening; knockout mouse; transgenic animal; ion channel disorder;  
XX fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
XX HERG; ss.

OS Homo sapiens.

XX WO200063434-A1.

XX 26-OCT-2000.

PF 14-APR-2000; 2000MO-US10004.

XX 15-APR-1999; 99US-0129404.

PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYIA ) UNIV YALE.

XX Abbott GW, Seidl F, Splawski I, Keating MT, Goldstein SAN;

XX WPI: 2000-672747/65.

DR P-PSDB; AAB29585.

PT Novel nucleic acids encoding MiRP1, MiRP2 and MiRP3, useful for

PT diagnosing and treating ion channel disorders, especially long QT

PT syndrome -

XX Claim 1; page 118-119; 132pp; English.

CC The invention relates to novel ion channel proteins related to  
CC KCNE1 (MiRP) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MiRP2; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MiRP3; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MiRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-KR), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents cDNA encoding human KCNE2 (MiRP1).  
XX

SO Sequence 732 BP; 221 A; 152 C; 157 G; 202 T; 0 other;

Query Match 100.0%; Score 372; DB 21; Length 732;  
Best Local Similarity 100.0%; Pred. No. 2.2e-102;

Matches 372; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGTCTACTTATTCATTTTCACACAGACGCTGGAGACGCTCCGAGAGATTATTAT 60  
DB 74 ATGTCTACTTATTCATTTTCACACAGACGCTGGAGACGCTCCGAGAGATTATTAT 133  
OY 61 ACTTATATGACAAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 120  
DB 134 ACTTATATGACAAATTTGGCGCCAGACACAAACAGCTGAGAGAGCCCTCCAGCCAAA 193  
OY 121 GTTATATGCTGAGAACTTCTACTATATCTCTGACTCATGTGATGATTTGAAATGTC 180  
DB 194 GTTATATGCTGAGAACTTCTACTATATCTCTGACTCATGTGATGATTTGAAATGTC 253  
OY 181 TCTTTCATCATCGTGGCCATCTGTGAGACAGCTGAAATCCAAAGACGGAACTCC 240  
DB 254 TCTTTCATCATCGTGGCCATCTGTGAGACAGCTGAAATCCAAAGACGGAACTCC 313  
OY 241 AATGACCCCTACACACAGTACATTTAGAGAGACTGGCAGAGAAAAGTCAAGAGCCAAATC 300  
DB 314 AATGACCCCTACACACAGTACATTTAGAGAGACTGGCAGAGAAAAGTCAAGAGCCAAATC 373  
OY 301 TTGAAATCTAGAAATGGAAGCCACCATTCATGAGAAACATTTGGCGCTGGTTCAAA 360  
DB 374 TTGAAATCTAGAAATGGAAGCCACCATTCATGAGAAACATTTGGCGCTGGTTCAAA 433  
OY 361 ATGTCCCTCTGA 372  
DB 434 ATGTCCCTCTGA 445

RESULT 9  
AAC64083  
ID AAC64083 standard; DNA; 732 BP.

XX AAC64083;  
AC  
XX  
DT 19-FEB-2001 (first entry)  
XX  
DE Human potassium channel protein KCNE2 (MIRP1) Q9E mutant DNA.  
XX  
XX Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KM Mink-related; long QT syndrome; cardiac arrhythmia;  
KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KM HERG; mutant; ds.  
XX  
XX Homo sapiens.  
OS Synthetic.  
XX  
PN MO200063434-A1.  
XX  
PD 26-OCT-2000.  
XX  
PF 14-APR-2000; 2000MO-US10004.  
XX  
PR 15-APR-1999; 99US-0129404.  
XX  
PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYA ) UNIV YALE.  
XX  
PI Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAN;  
XX WPI; 2000-672747/65.  
DR P-PSDB; AAB29593.  
XX  
XX Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
PT diagnosing and treating ion channel disorders, especially long QT  
PT syndrome -  
XX  
XX Claim 56; Page -: 132pp; English.  
XX  
XX The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
CC potassium channels (I-Kr), mutations in which are associated with long  
CC QT syndrome. The invention also relates to methods of diagnosing long QT  
CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
CC nonhuman animals comprising a heterologous ion channel protein gene  
CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
CC DNA, and methods of and screening drugs for treating long QT syndrome  
CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
CC acids, and proteins may be used for diagnosing or treating ion channel  
CC disorders, especially long QT syndrome. Transgenic animals comprising  
CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
CC The present sequence represents DNA encoding a mutant human KCNE2  
CC (MIRP1) specifically claimed for use in diagnostic and drug screening  
CC methods of the invention.  
CC Note: The present sequence is not shown in the specification, but is  
CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
CC 118-119.  
XX  
XX Sequence 732 BP; 221 A; 151 C; 158 G; 202 T; 0 other;  
SO  
Query Match 99.6%; Score 370.4; DB 21; Length 732;  
Best Local Similarity 99.7%; Pred. No. 6.7e-102;  
Matches 371; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
QY 1 ATGTCTACTTATCCAAATTCACACAGACGCTGGAAGAGCTCTCCGAAGATTTTATT 60  
DB 74 ATGTCTACTTATCCAAATTCACACAGACGCTGGAAGAGCTCTCCGAAGATTTTATT 133

QY 61 ACTTATATGACAAATTTGGCGCCAGAACACAAACAGCTGAGCAAGAGCCCTCCAAAGCCAAA 120  
DB 134 ACTTATATGACAAATTTGGCGCCAGAACACAAACAGCTGAGCAAGAGCCCTCCAAAGCCAAA 193  
QY 121 GTTGATGTGAGAACTTCTACTATGTCACTCTGTACCTCATGTGATGATTTGAATGTTTC 180  
DB 194 GTTGATGTGAGAACTTCTACTATGTCACTCTGTACCTCATGTGATGATTTGAATGTTTC 253  
QY 181 TCTTTCATCATGTGTGGCCATCTGTGTGAGCACTGTGAATTCACAGACGCGGAACATCTCC 240  
DB 254 TCTTTCATCATGTGTGGCCATCTGTGTGAGCACTGTGAATTCACAGACGCGGAACATCTCC 313  
QY 241 AATGACCCCTTACACAGATCATGTGTAGAGCACTGTGAGCAAGAAAGTACAAAGCCAAATTC 300  
DB 314 AATGACCCCTTACACAGATCATGTGTAGAGCACTGTGAGCAAGAAAGTACAAAGCCAAATTC 373  
QY 301 TTGAATCTAGAAAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGGGTTCCAAA 360  
DB 374 TTGAATCTAGAAAGAAATCGAAGGCCACATCCATGAGAAATTTGGTGGGTTCCAAA 433  
QY 361 ATGTCCCTCTGA 372  
DB 434 ATGTCCCTCTGA 445  
RESULT 10  
AAC64084  
ID AAC64084 standard; DNA: 732 BP.  
XX  
XX AAC64084;  
XX  
DT 19-FEB-2001 (first entry)  
XX  
XX Human potassium channel protein KCNE2 (MIRP1) M54T mutant DNA.  
DE  
XX Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
KM Mink-related; long QT syndrome; cardiac arrhythmia;  
KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
KM HERG; mutant; ds.  
XX  
XX Homo sapiens.  
OS Synthetic.  
XX  
PN MO200063434-A1.  
XX  
PD 26-OCT-2000.  
XX  
PF 14-APR-2000; 2000MO-US10004.  
XX  
PR 15-APR-1999; 99US-0129404.  
XX  
PA (UTAH ) UNIV UTAH RES FOUND.  
PA (UYA ) UNIV YALE.  
XX  
PI Abbott GW, Sesti F, Splawski I, Keating MT, Goldstein SAN;  
XX WPI; 2000-672747/65.  
DR P-PSDB; AAB29594.  
XX  
XX Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
PT diagnosing and treating ion channel disorders, especially long QT  
PT syndrome -  
XX  
XX Claim 56; Page -: 132pp; English.  
XX  
XX The invention relates to novel ion channel proteins related to  
CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
CC respectively). The cDNAs encoding these proteins are given in AAC64071-



Db 314 AATGACCCCTACACAGTACATTGTAGAGACTGGCAGGAAAAAGTACAAGACCAATC 373  
 QY 301 TTGAATCTAGAAATCGAAGGCCACCATTCATGAGAACTTGTGCGGTGCTCAAA 360  
 Db 374 TTGAATCTAGAAATCGAAGGCCACCATTCATGAGAACTTGTGCGGTGCTCAAA 433  
 QY 361 ATGTCCCTCTGA 372  
 Db 434 ATGTCCCTCTGA 445  
 RESULT 12  
 AAC64086  
 ID AAC64086 standard; DNA; 732 BP.  
 AC AAC64086;  
 DT 19-FEB-2001 (first entry)  
 XX  
 DE Human potassium channel protein KCNE2 (MIRP1) T9A mutant DNA.  
 XX  
 KM Human; KCNE2; MIRP1; potassium channel protein; KCNE1-related;  
 KM Mink-related; long QT syndrome; cardiac arrhythmia;  
 KM drug screening; knockout mouse; transgenic animal; ion channel disorder;  
 KM fast delayed rectifier potassium channel; anti-KCNE2 antibody;  
 KM HERG; mutant; ds.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 OS  
 PN WO200063434-A1.  
 XX  
 PD 26-OCT-2000.  
 XX  
 PE 14-APR-2000; 2000WO-US10004.  
 PR 15-APR-1999; 99US-0129404.  
 PR  
 PA (UTAH ) UNIV UTAH RES FOUND.  
 PA (UYVA ) UNIV YALE.  
 PI Abbott GW, Seesti F, Splawski I, Keating MT, Goldstein SAN;  
 DR MPI; 2000-672747/65.  
 DR P-PDB; AAB29596.  
 XX  
 PT Novel nucleic acids encoding MIRP1, MIRP2 and MIRP3, useful for  
 PT diagnosing and treating ion channel disorders, especially long QT  
 PT syndrome -  
 XX  
 PS Claim 56; Page -: 132pp; English.  
 XX  
 CC The invention relates to novel ion channel proteins related to  
 CC KCNE1 (Mink) and to nucleic acids encoding them. The proteins of  
 CC the invention are human and rat KCNE2 (MIRP1; AAB29585 and AAB29586,  
 CC respectively); human and mouse KCNE3 (MIRP2; AAB29587 and AAB29588,  
 CC respectively); and human and mouse KCNE4 (MIRP3; AAB29589 and AAB29590,  
 CC respectively). The cDNAs encoding these proteins are given in AAC64071-  
 CC AAC64076. KCNE2, along with HERG, forms cardiac fast delayed rectifier  
 CC potassium channels (I-KR), mutations in which are associated with long  
 CC QT syndrome. The invention also relates to methods of diagnosing long QT  
 CC syndrome using the KCNE2, KCNE3 or KCNE4 genes, a knockout mouse with a  
 CC disruption in an endogenous KCNE2, KCNE3 or KCNE4 gene, transgenic  
 CC nonhuman animals comprising a heterologous ion channel protein gene  
 CC of the invention, a transgenic animal comprising human KCNE2 and HERG  
 CC DNA, and methods of and screening drugs for treating long QT syndrome  
 CC using KCNE2 proteins (including mutants), nucleic acids encoding them  
 CC and antibodies against KCNE2 proteins. The methods, antibodies, nucleic  
 CC acids, and proteins may be used for diagnosing or treating ion channel  
 CC disorders, especially long QT syndrome. Transgenic animals comprising  
 CC KCNE2 and HERG are useful for testing anti-long QT syndrome drugs.  
 CC The present sequence represents DNA encoding a mutant human KCNE2  
 CC (MIRP1) specifically claimed for use in diagnostic and drug screening

CC methods of the invention.  
 CC Note: The present sequence is not shown in the specification, but is  
 CC derived from the wild-type human KCNE2 cDNA sequence shown on page  
 CC 118-119.  
 XX  
 SQ Sequence 732 BP; 220 A; 152 C; 158 G; 202 T; 0 other;  
 Query Match 99.6%; Score 370.4; DB 21; Length 732;  
 Best Local Similarity 99.7%; Pred. No. 6.7e-102;  
 Matches 371; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 ATGTCTACTTTATCCAAATTCACAGACGCTGGAAGAGCTTTCGGAAGATTTTATTT 60  
 Db 74 ATGTCTACTTTATCCAAATTCGACAGACGCTGGAAGAGCTTTCGGAAGATTTTATTT 133  
 QY 61 ACTTATATGACAAATTTGGCGCCAGAAACACACAGCTGAGCAAGAGCCCTCCAGCCAAA 120  
 Db 134 ACTTATATGACAAATTTGGCGCCAGAAACACACAGCTGAGCAAGAGCCCTCCAGCCAAA 193  
 QY 121 GTTGATGCTGAGAACTTCTACTATGTCATGTCATGTCATGTCATGTCATGTCATGTCAT 180  
 Db 194 GTTGATGCTGAGAACTTCTACTATGTCATGTCATGTCATGTCATGTCATGTCATGTCAT 253  
 QY 181 TCTTTCATCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCAT 240  
 Db 254 TCTTTCATCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCATGTCAT 313  
 QY 241 AATGACCCCTACACAGTACATTGTAGAGAGCTGGCAGGAAAGTACAGACCAATTC 300  
 Db 314 AATGACCCCTACACAGTACATTGTAGAGAGCTGGCAGGAAAGTACAGACCAATTC 373  
 QY 301 TTGAATCTAGAAATCGAAGGCCACCATTCATGAGAACTTGTGCGGTGCTCAAA 360  
 Db 374 TTGAATCTAGAAATCGAAGGCCACCATTCATGAGAACTTGTGCGGTGCTCAAA 433  
 QY 361 ATGTCCCTCTGA 372  
 Db 434 ATGTCCCTCTGA 445  
 RESULT 13  
 ABA49938  
 ID ABA49938 standard; DNA; 312 BP.  
 AC ABA49938;  
 DT 01-FEB-2002 (first entry)  
 XX  
 DE Human breast cell single exon nucleic acid probe #8633.  
 XX  
 KM Human; microarray; single exon probe; gene expression; breast;  
 KM disease; cancer; ss.  
 OS Homo sapiens.  
 OS  
 PN WO200157271-A2.  
 XX  
 PD 09-AUG-2001.  
 XX  
 PE 30-JAN-2001; 2001WO-US00662.  
 PR 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024263.  
 XX  
 PA (MOLE-) MOLECULAR DYNAMICS INC.  
 PA Penn SG, Hanzel DK, Chen W, Rank DR;  
 PI

DR WPI: 2001-496933/54.  
XX  
PT New spatially-addressable set of single exon nucleic acid probes,  
PT useful for measuring gene expression in sample derived from human  
PT breast, comprises number of single exon nucleic acid probes -  
XX  
PS  
XX Claim 4: SEQ ID NO 8633: 327bp + sequence listing; English.  
XX  
CC The invention relates to a spatially-addressable set of single exon  
CC nucleic acid probes for measuring gene expression in a sample derived  
CC from human breast and BT 474 cells. The method involves contacting  
CC the probes with a collection of detectably labelled nucleic acids  
CC derived from mRNA of human breast, and then measuring the label  
CC bound to each probe of the microarray. The probes are useful for  
CC verifying the expression of regions of genomic DNA predicted to  
CC encode proteins. They are useful for gene discovery, and for  
CC determining predisposition and/or prognosing breast disease. Gene  
CC expression analysis is useful for assessing the toxicity of chemical  
CC agents on cells. The microarray of this invention presents a far greater  
CC diversity of probes for measuring gene expression, with far less bias  
CC than expressed sequence tag microarrays. The method is suitable for  
CC rapid production of functional information from genomic sequence. The  
CC present sequence is a single exon nucleic acid probe of the invention.  
CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pcl\_sequences.  
XX  
SQ Sequence 312 BP: 94 A; 73 C; 71 G; 74 T; 0 other;  
Query Match 83.9%; Score 312; DB 22; Length 312;  
Best Local Similarity 100.0%; Pred. No. 2e-84; Indels 0; Gaps 0;  
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 54 TTTTATTACTTATATGACATTTGGCGCCAGAACACAGCTGAGCAGAGGCCCTCCA 113  
DB 1 TTTTATTACTTATATGACATTTGGCGCCAGAACACAGCTGAGCAGAGGCCCTCCA 60  
OY 114 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTGATCTCATTTGGTGGATTGG 173  
DB 61 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTGATCTCATTTGGTGGATTGG 120  
OY 174 AATGTCCTCTTTCATATCGTGGCCATCTGCTGAGCAGCTGTGAATCCAAAGAGCGGA 233  
DB 121 AATGTCCTCTTTCATATCGTGGCCATCTGCTGAGCAGCTGTGAATCCAAAGAGCGGA 180  
OY 234 ACACCTCAATGAGCCCTTACCAACAGTACATTGTAGAGAGCTGGCAGGAAAAGTACAAGAG 293  
DB 181 ACACCTCAATGAGCCCTTACCAACAGTACATTGTAGAGAGCTGGCAGGAAAAGTACAAGAG 240  
OY 294 CCAAAATCTGAATCTAGAAAGATCGAAGCCACATCCATGAGAACTTGGTGGCTGG 353  
DB 241 CCAAAATCTGAATCTAGAAAGATCGAAGCCACATCCATGAGAACTTGGTGGCTGG 300  
OY 354 GTTCAAAATGTC 365  
DB 301 GTTCAAAATGTC 312  
RESULT 14  
ID ABA67856 standard; DNA: 312 BP.  
XX ABA67856:  
XX  
XX 01-FEB-2002 (first entry)  
XX  
XX Human foetal liver single exon nucleic acid probe #16161.  
XX  
XX Human: foetal liver; gene expression; single exon nucleic acid probe: ss.  
XX  
XX Homo sapiens.  
XX  
XX MO200157277-A2.  
XX  
XX

XX  
PD 09-AUG-2001.  
XX  
PF 30-JAN-2001: 2001MO-US00669.  
XX  
PR 04-FEB-2000: 2000US-0180312.  
PR 26-MAY-2000: 2000US-0207456.  
PR 30-JUN-2000: 2000US-0608408.  
PR 03-AUG-2000: 2000US-0632365.  
PR 21-SEP-2000: 2000US-0234687.  
PR 27-SEP-2000: 2000US-0236359.  
PR 04-OCT-2000: 2000US-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
XX WPI: 2001-483447/52.  
XX  
XX Human genome-derived single exon nucleic acid probes useful for  
XX analyzing gene expression in human foetal liver -  
XX  
PS Claim 4: SEQ ID NO 16161: 639pp + sequence listing; English.  
XX  
XX The invention relates to a single exon nucleic acid probe for  
XX measuring human gene expression in a sample derived from human foetal  
XX liver. The single exon nucleic acid probes may be used for predicting,  
XX measuring and displaying gene expression in samples derived from human  
XX foetal liver. The present sequence is a single exon nucleic acid  
XX probe of the invention.  
XX Note: The sequence data for this patent did not form part of the  
XX printed specification, but was obtained in electronic format directly  
XX from WIPO at ftp.wipo.int/pub/published\_pcl\_sequences.  
XX  
SQ Sequence 312 BP: 94 A; 73 C; 71 G; 74 T; 0 other;  
Query Match 83.9%; Score 312; DB 22; Length 312;  
Best Local Similarity 100.0%; Pred. No. 2e-84; Indels 0; Gaps 0;  
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 54 TTTTATTACTTATATGACATTTGGCGCCAGAACACAGCTGAGCAGAGGCCCTCCA 113  
DB 1 TTTTATTACTTATATGACATTTGGCGCCAGAACACAGCTGAGCAGAGGCCCTCCA 60  
OY 114 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTGATCTCATTTGGTGGATTGG 173  
DB 61 AGCCAAAGTTGATGCTGAGAACTTCTACTATGTATCTCTGATCTCATTTGGTGGATTGG 120  
OY 174 AATGTCCTCTTTCATATCGTGGCCATCTGCTGAGCAGCTGTGAATCCAAAGAGCGGA 233  
DB 121 AATGTCCTCTTTCATATCGTGGCCATCTGCTGAGCAGCTGTGAATCCAAAGAGCGGA 180  
OY 234 ACACCTCAATGAGCCCTTACCAACAGTACATTGTAGAGAGCTGGCAGGAAAAGTACAAGAG 293  
DB 181 ACACCTCAATGAGCCCTTACCAACAGTACATTGTAGAGAGCTGGCAGGAAAAGTACAAGAG 240  
OY 294 CCAAAATCTGAATCTAGAAAGATCGAAGCCACATCCATGAGAACTTGGTGGCTGG 353  
DB 241 CCAAAATCTGAATCTAGAAAGATCGAAGCCACATCCATGAGAACTTGGTGGCTGG 300  
OY 354 GTTCAAAATGTC 365  
DB 301 GTTCAAAATGTC 312  
RESULT 15  
ID ABA34913 standard; DNA: 312 BP.  
XX ABA34913:  
XX  
XX 23-JAN-2002 (first entry)  
XX  
XX

DE Probe #13379 for gene expression analysis in human heart cell sample.  
XX  
XX Human: gene expression; heart; microarray; vascular system; probe;  
KM cardiovascular disease; hypertension; cardiac arrhythmia;  
KW congenital heart disease; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX MO200157274-A2.  
XX  
XX  
XX PD 09-AUG-2001.  
XX  
XX  
XX PF 30-JAN-2001; 2001WO-US00666.  
XX  
XX PR 04-FEB-2000; 2000US-0180312.  
XX PR 26-MAY-2000; 2000US-0207456.  
XX PR 30-JUN-2000; 2000US-0608408.  
XX PR 03-AUG-2000; 2000US-0632366.  
XX PR 21-SEP-2000; 2000US-0234687.  
XX PR 27-SEP-2000; 2000US-0236359.  
XX PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX  
XX PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
XX PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
XX DR WPI; 2001-488899/53.  
XX  
XX PT Single exon nucleic acid probes for analyzing gene expression in human  
XX hearts -  
XX  
XX PS Claim 4; SEQ ID NO 13379; 530bp; English.  
XX

CC The present invention relates to single exon nucleic acid probes for  
CC measuring human gene expression in a sample derived from human heart. The  
CC present sequence is one such probe. The probes may be used for  
CC predicting, measuring and displaying gene expression in samples derived  
CC from the human heart via microarrays. By measuring gene expression, the  
CC probes are useful for predicting, diagnosing, grading, staging,  
CC monitoring and prognosing diseases of the human heart and vascular system  
CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and  
CC congenital heart disease.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX  
XX SQ Sequence 312 BP; 94 A; 73 C; 71 G; 74 T; 0 other;

Query Match 83.9%; Score 312; DB 22; Length 312;  
Best local Similarity 100.0%; Pred. No. 2e-84;  
Matches 312; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 54 TTTTATTACTTATATGACATTTGGCCGACGACACACAGCTGAGCAAGAGGCGCTCA 113  
DB 1 TTTTATTACTTATATGACATTTGGCCGACGACACACAGCTGAGCAAGAGGCGCTCA 60  
OY 114 AGCCAAAGTTGATGCTGAGAACTTCTACATCTGATCCCTGATCCATGATGATTTGG 173  
DB 61 AGCCAAAGTTGATGCTGAGAACTTCTACATCTGATCCCTGATGATGATTTGG 120  
OY 174 AATGTTCTCTTCATCATGCTGGCCATCTGTGTGAGCAGCTGTAATCCAAAGAGCGGA 233  
DB 121 AATGTTCTCTTCATCATGCTGGCCATCTGTGTGAGCAGCTGTAATCCAAAGAGCGGA 180  
OY 234 ACACTCCATGACCTTACCCAGCAGTACATTTAGAGAGCTGGCAGAAAAGTACAAGAG 293  
DB 181 ACACTCCATGACCTTACCCAGCAGTACATTTAGAGAGCTGGCAGAAAAGTACAAGAG 240  
OY 294 CCAATCTGAATCTAGAGAAATCGAAGGCCATCCATCCATGAGAAATGCTGGCGCTGG 353  
DB 241 CCAATCTGAATCTAGAGAAATCGAAGGCCATCCATCCATGAGAAATGCTGGCGCTGG 300  
OY 354 GTTCAAAATGTC 365

DB 301 GTTCAAAATGTC 312  
|||||

Search completed: October 24, 2002, 13:19:51  
Job time : 201 secs





GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:16:15 : Search time 43 seconds  
(without alignments)  
2125.015 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_445

Perfect score: 372  
Sequence: 1 atgtctattatcccaatt.....gttcaaaatgtcccccctga 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 383533 seqs, 122816752 residues

Total number of hits satisfying chosen parameters: 767066

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued\_Patents\_NA:\*

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- 3: /cgn2\_6/ptodata/2/1na/6A\_COMB.seq:\*
- 4: /cgn2\_6/ptodata/2/1na/6B\_COMB.seq:\*
- 5: /cgn2\_6/ptodata/2/1na/PC105\_COMB.seq:\*
- 6: /cgn2\_6/ptodata/2/1na/backfiles1.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	53.2	14.3	398	US-08-118-101A-5	Sequence 5, Appl1
2	53.2	14.3	1703	US-09-135-021-77	Sequence 77, Appl1
3	53.2	14.3	1703	US-09-135-020-3	Sequence 3, Appl1
4	53.2	14.3	1703	US-09-135-010A-3	Sequence 3, Appl1
5	36.6	9.8	2652	US-08-318-831-1	Sequence 1, Appl1
6	36	9.7	1380	US-08-110-286A-1	Sequence 1, Appl1
7	33.6	9.0	645	US-09-069-896-2	Sequence 2, Appl1
8	33	8.9	606	US-09-328-111-133	Sequence 133, App
9	33	8.9	2912	US-08-998-416-303	Sequence 3, Appl1
10	32.4	8.7	837	US-08-998-416-303	Sequence 303, App
11	30.6	8.2	6822	US-09-426-998-3	Sequence 3, Appl1
12	30.6	8.2	7741	US-08-724-394A-18	Sequence 18, Appl1
13	29.8	8.0	2266	US-08-724-394A-18	Sequence 20, Appl1
14	29.8	8.0	246240	US-08-724-394A-20	Sequence 21, Appl1
15	29.8	8.0	246240	US-08-724-394A-21	Sequence 22, Appl1
16	29.8	8.0	246240	US-08-724-394A-22	Sequence 23, Appl1
17	29.6	8.0	462	US-08-863-813A-33	Patent No. 551630
18	29.6	8.0	3273	US-08-475-886-5	Sequence 5, Appl1
19	29.6	8.0	7486	US-08-397-232-3	Sequence 3, Appl1
20	29.6	8.0	7486	US-08-475-886-3	Sequence 1, Appl1
21	29.6	8.0	7488	US-08-475-886-1	Sequence 1, Appl1
22	29.6	8.0	7493	US-08-397-232-1	Sequence 1, Appl1
23	29.6	8.0	7493	US-09-171-387-1	Sequence 1, Appl1
24	29.6	8.0	7493	US-08-975-762-47	Sequence 47, Appl1
25	29.4	7.9	3947	US-09-295-028-47	Sequence 47, Appl1
26	29.4	7.9	3947	US-09-106-582-47	Sequence 47, Appl1
27	29.4	7.9	3947		

C 28	29.2	7.8	352	4	US-09-439-313-421	Sequence 421, App
C 29	29.2	7.8	630	4	US-08-646-695-13	Sequence 13, Appl
C 30	29.2	7.8	630	4	US-08-646-695-14	Sequence 14, Appl
C 31	29.2	7.8	630	5	PCT-US96-06053-13	Sequence 13, Appl
C 32	29.2	7.8	630	5	PCT-US96-06053-14	Sequence 14, Appl
C 33	29.2	7.8	2380	1	US-08-572-951-3	Sequence 3, Appl1
C 34	29.2	7.8	14311	4	US-08-646-695-1	Sequence 1, Appl1
C 35	29.2	7.8	14311	4	US-08-646-695-7	Sequence 7, Appl1
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C 38	29	7.8	744	3	US-08-969-644-17	Sequence 17, Appl
C 39	29	7.8	744	3	US-08-444-189-17	Sequence 17, Appl
C 40	29	7.8	744	4	US-08-468-544-17	Sequence 17, Appl
C 41	29	7.8	1983	1	US-08-073-799C-9	Sequence 9, Appl1
C 42	29	7.8	7502	3	US-08-969-644-6	Sequence 6, Appl1
C 43	29	7.8	7502	3	US-08-444-189-6	Sequence 6, Appl1
C 44	29	7.8	7502	4	US-08-468-544-6	Sequence 6, Appl1
C 45	28.8	7.7	2168	2	US-08-633-879C-1	Sequence 1, Appl1

## ALIGNMENTS

RESULT 1  
US-08-118-101A-5  
Sequence 5, Application US/08118101A  
Patent No. 5620892  
GENERAL INFORMATION:  
APPLICANT: Kutz, Stephen E.  
APPLICANT: Knickerbocker, Aron M.  
APPLICANT: McCullough, John R.  
TITLE OF INVENTION: A STRAIN OF SACCAROMYCES CEREVISIAE  
TITLE OF INVENTION: EXPRESSING THE GENE ENCODING POTASSIUM TRANSPORTER MINK  
NUMBER OF SEQUENCES: 16  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Burton Rodney  
STREET: P.O. Box 4000  
CITY: Princeton  
STATE: New Jersey  
COUNTRY: U.S.A.  
ZIP: 08543-4000  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/118,101A  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Gaul, Timothy J.  
REGISTRATION NUMBER: 33,111  
REFERENCE/DOCKET NUMBER: DC27  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (609) 252-5901  
TELEFAX: (609) 252-4526  
INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 398 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..398  
US-08-118-101A-5  
Query Match 14.3%, Score 53.2, DB 1; Length 398;  
Best Local Similarity 63.6%, Pred. No. 1.3e+08;  
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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Db 141 CCTTACGTCTCTCATGTGTACTGTGGGATTTCTTGCGTTTCTTACCCCTGGGCTCATGTGTGAG 200

QY 210 CACGTGAATTCAGAGAGAGGGGACACTCCATGACCCTTACCACAGTACATTG---T 266

Db 201 CTACATCCGCTCCCAAGAGCTGGGAGCACTGGAAGACCATTCAACGCTTCATACGAGTIC 260

QY 267 AGAGACATGGCAGGAAAAGTACAGACCAATTC 300

Db 261 CGATGCTTGGCAGAGAGACGACCAAGGCTTATGTC 294

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RESULT 2
US-09-135-021-77
? Sequence 77, Application US/09135021A
? Patent No. 6130104
? GENERAL INFORMATION:
? APPLICANT: Splawski, Igor
? APPLICANT: Keating, Mark T.
? TITLE OF INVENTION: A HOMOTYGOUS MUTATION IN KYLOT1 WHICH CAUSES JERVELL
? TITLE OF INVENTION: AND LANGE-NIELSEN SYNDROME
? FILE REFERENCE: 2323-128
? CURRENT APPLICATION NUMBER: US/09/135,021A
? CURRENT FILING DATE: 1998-08-17
? EARLIER APPLICATION NUMBER: 08/874,655
? EARLIER FILING DATE: 1997-06-13
? EARLIER APPLICATION NUMBER: 60/094,477
? EARLIER FILING DATE: 1998-07-29
? NUMBER OF SEQ ID NOS: 80
? SOFTWARE: PatentIn Ver. 2.0
? SEQ ID NO 77
? LENGTH: 1703
? TYPE: DNA
? ORGANISM: Homo sapiens
? FEATURE:
? NAME/KEY: CDS
? LOCATION: (193)..(579)
US-09-135-021-77

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Query Match: 14.3% Score 53.2; DB 3; Length 1703;
Best Local Similarity 63.6%; Pred. No. 3e-08;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1

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DB 324 CCTGTACGTCCTCATGTGATGATTTGATGATGTCTTTCATTCATCGTGGCCATCCTGTGTAG 383

QY 210 CACTGTGAATTCACACAGACGGGAACACTCCCAATACCCCTCACACAGTACATG---T 266
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DB 384 CTACATCTCGCTCCACAGAACCTGTGACACACTCGAACACCCATTCACATTCATCGATGC 443

QY 267 AGAGGACTGGCAGAGAAAGTACAGAGCCAAATC 300
      || | ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
DB 444 CGATGCTGGCAGAGAACAGACAGAGCCATGTC 477

RESULT 3
US-09-135-020-3
: Sequence 3, Application US/09135020
: Patent No. 6274332
: GENERAL INFORMATION:
: APPLICANT: Keating, Mark T.
: APPLICANT: Sanguinetti, Michael C.
: APPLICANT: Splavski, Igor
: TITLE OF INVENTION: MOTATIONS IN THE KNEEL GENE ENCODING HUMAN MINK WHICH
: TITLE OF INVENTION: CAUSE ARRHYNHMA SUSCEPTIBILITY THEREBY ESTABLISHING
: FILE REFERENCE: 2323-131
: CURRENT APPLICATION NUMBER: US/09/135,020
: CURRENT FILING DATE: 1998-08-17
: EARLIER APPLICATION NUMBER: 08/921,068
: EARLIER FILING DATE: 1997-08-29

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	EARLIER APPLICATION NUMBER: 08/739,383	
	EARLIER FILING DATE: 1996-10-29	
	EARLIER APPLICATION NUMBER: 60/019,014	
	EARLIER FILING DATE: 1995-12-22	
	EARLIER APPLICATION NUMBER: 60/094,477	
	EARLIER FILING DATE: 1998-07-29	
	NUMBER OF SEQ ID NOS: 114	
	SOFTWARE: PatentIn Ver. 2.0	
	SEQ ID NO 3	
	LENGTH: 1703	
	TYPE: DNA	
	ORGANISM: Homo sapiens	
	FEATURE:	
	NAME/KEY: CDS	
	LOCATION: (193)..(579)	
	US-09-135-020-3	
Query Match	14.3%; Score 53.2; DB 4; Length 1703;	
Best Local Similarity	63.6%; Pred. No 3e-08;	
Matches 98; Conservative	0; Mismatches 53; Indels 3; Gaps 1.	
OY	150 CCTGTACCTCATGTGATGATTTGAAATGTTCTCTTTCATCATCGTGGCCATCCTGTGTAG	209
Db	334 CCTTCACGTCCTCCCAAGGAGTACGTGGATTTCTCGCTTTCACCCGTGGGATCATGCTGTAG	383
OY	210 CACGTGAAATCCAGAGACGGGAACTCCAAATGACCCCTTACCCACAGATACATTG---T	266
Db	384 CTACATCGCGCTCCAGGAAGCTGGAGACATCTCGAAGACCCATTCACAAAGTCTACATCGAGATC	443
OY	267 AGAGGACTGGCAGAGAAAAGTACAAAGCCAAATC	300
Db	444 CGATGCTTGGCCAGGAAGGACCAAGGCTATGTC	477

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RESULT 4
US-09-135-010A-3
; Sequence 3, Application US/09135010A
; Patent No. 6277978
; GENERAL INFORMATION:
; APPLICANT: Keating, Mark T.
; APPLICANT: Sanguinetti, Michael C.
; APPLICANT: Curran, Mark E.
; APPLICANT: Landes, Gregory M.
; APPLICANT: Connors, Timothy D.
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: KVL0T1 - A LONG QT SYNDROME GENE
; FILE REFERENCE: 2323-133
; CURRENT APPLICATION NUMBER: US/09/135,010A
; CURRENT FILING DATE: 1998-08-17
; PRIOR APPLICATION NUMBER: 60/094,477
; PRIOR FILING DATE: 1998-07-29
; PRIOR APPLICATION NUMBER: 08/921,068
; PRIOR FILING DATE: 1997-08-29
; PRIOR APPLICATION NUMBER: 08/739,383
; PRIOR FILING DATE: 1996-10-29
; PRIOR APPLICATION NUMBER: 60/019,014
; PRIOR FILING DATE: 1995-12-22
; NUMBER OF SEQ ID NOS: 116
; SOFTWARE: PatentIn Ver.. 2.0
SEQ ID NO 3
; LENGTH: 1703
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (193)..(579)
; US-09-135-010A-3

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Query Match	14.3%	Score 53.2;	DB 4;	Length 1703;
Best Local Similarity	63.6%	Pred. No. 3e-08;		
Matches 98; Conservative	0;	Mismatches 53;	Indels 3;	Gaps 1;

malchines /6; conservative 0; mish

Page 3







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1  APPLICANT: Feder, John N.
2  APPLICANT: Krommal, Gregory S.
3  APPLICANT: Laufer, Peter M.
4  APPLICANT: Ruddy, David A.
5  APPLICANT: Thomas, Winston
6  APPLICANT: Tsuchihashi, Zenta
7  APPLICANT: Wolff, Roger K.
8  TITLE OF INVENTION: Megabase Transcript Map: No. 5872237el
9  TITLE OF INVENTION: Sequences and Antibodies Thereeto
10 NUMBER OF SEQUENCES: 31
11
12 CORRESPONDENCE ADDRESSES:
13 ADDRESSEE: TOWNSEND and TOWNSEND and CREW LLP
14 STREET: Two Embarcadero Center, 8th Floor
15 CITY: San Francisco
16 STATE: CA
17 COUNTRY: USA
18 ZIP: 94111-3834
19
20 COMPUTER READABLE FORM:
21 MEDIUM TYPE: Floppy disk
22 COMPUTER: IBM PC compatible
23 OPERATING SYSTEM: PC-DOS/MS-DOS
24 SOFTWARE: PatentIn Release #1.0, Version #1.30
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26 CURRENT APPLICATION DATA:
27 APPLICATION NUMBER: US/08/724,394A
28 FILING DATE: 01-OCT-1996
29 CLASSIFICATION: 536
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31 ATTORNEY/AGENT INFORMATION:
32 NAME: Fitts, Renee A.
33 REGISTRATION NUMBER: 35,136
34 REFERENCE/DOCKET NUMBER: 017957-000100
35 TELECOMMUNICATION INFORMATION:
36 TELEPHONE: 415-576-0200
37 TELEFAX: 415-576-0300
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39 INFORMATION FOR SEQ ID NO: 21:
40 SEQUENCE CHARACTERISTICS:
41 LENGTH: 246240 base pairs
42 TYPE: nucleic acid
43 STRANDEDNESS: not relevant
44 TOPOLOGY: not relevant
45 MOLECULE TYPE: cDNA
46 FEATURE:
47 NAME/KEY: misc feature
48 LOCATION: 1..246240
49 OTHER INFORMATION: /note= "HLA-H CONTIG"
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Best Local Similarity	60.5%	Pred. No. 50		
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QY	212	CTGTGAATCCCAAGACGCG	232	
Db	29076	TTTGAGAAAGTCATGCTAATAAG	29096	

Search completed: October 24, 2002, 14:20:05  
Job time : 147 secs





GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: October 24, 2002, 13:15:40 : Search time 1605 Seconds

(without alignments)  
3128.266 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_445

Perfect score: 372  
Sequence: 1 atgtctattctatccattt.....gttcaaaatgtcccccctga 372

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 674847542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-Processing: Minimum Match 0%

Listing first 45 summaries

Database :

EST:\*  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estinu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_gss:\*  
13: em\_gss\_hum:\*  
14: em\_gss\_inv:\*  
15: em\_gss\_pln:\*  
16: em\_gss\_vrt:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	357.8	96.2	803	10	BG208163 RST7654
2	336	90.3	410	9	AI962650
3	336	90.3	429	9	AI654552
4	331.8	89.2	391	9	AI339609
5	338	88.2	372	9	AI246239
6	269.6	72.5	1691	11	AK008619
7	260.8	70.1	470	10	DB5797
8	245	65.9	1003	10	BG261965
9	177.8	47.8	756	10	BG21966
10	168.2	45.2	341	10	BG938225
11	160	43.0	188	9	AA633404
12	119.4	32.1	121	9	AA935321
13	99	26.6	311	9	AM869303
14	90	24.2	314	9	BB564873
15	57.6	15.5	272	9	BB574249
16	57.2	15.4	358	10	BE486735
17	53.2	14.3	716	10	BI459541

18	52.8	14.2	464	9	AA67912
19	52.8	14.2	603	9	AT956381
20	52.8	14.2	674	9	BB613272
21	52.8	14.2	754	11	AK008938
22	51.8	13.9	311	9	AM869303
23	51	13.7	424	10	BM389584
24	50.8	13.7	270	9	BB595946
25	50.8	13.7	869	10	BF540248
26	45	12.1	986	12	CNS076KL
27	42.2	11.3	484	10	BU059218
28	42.2	11.3	687	10	BU098508
29	42	11.3	597	10	BU038615
30	40.4	10.9	589	10	BU094875
31	40.4	10.9	629	10	BU095114
32	38.8	10.7	309	10	BU059046
33	38.4	10.3	634	9	AM187570
34	38.4	10.3	694	10	BF479357
35	38.4	10.3	697	10	BM301804
36	37.8	10.2	455	10	BG554741
37	37.2	10.0	539	10	BI450048
38	36.2	9.7	431	9	BB789522
39	36.2	9.7	772	10	BE273157
40	36	9.7	1393	10	BF131615
41	35.8	9.6	538	9	AV883212
42	35.8	9.6	673	9	AV889968
43	35.8	9.6	675	9	AV676222
44	35.8	9.6	709	9	AV858687
45	35.6	9.6	578	9	AV391303

## ALIGNMENTS

RESULT 1	BG208163/c	803 bp	MRNA	linear	EST 21-APR-2001
LOCUS	RST7654	Athersys RAGE Library	Homo sapiens	cDNA	RNA sequence.
DEFINITION	BG208163				
ACCESSION	BG208163				
VERSION	BG208163.1	GI:13729850			
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 803)				
AUTHORS	Harrington,J.J., Sherf,B., Rundlett,S., Jackson,P.D., Perry,R., Cain,S., Leventhal,C., Thornton,M., Ramachandran,R., Whittington,J., Lerner,L., Costanzo,D., McElligott,K., Booser,S., Maye,R., Smith,E., Veloso,N., Klika,A., Hess,J., Cothren,K., Lo,K., Offenbacher,J., Danzig,J. and Ducar,M.				
TITLE	Creation of genome-wide protein expression libraries using random activation of gene expression				
JOURNAL	Nat. Biotechnol. 19 (5), 440-445 (2001)				
MEDLINE	21227151				
COMMENT	Contact: Scott J. Cain				
	Athersys, Inc.				
	3201 Carnegie Ave, Cleveland, OH 44115, USA				
	Tel: 216 431 9900				
	Fax: 216 361 9596				
	Email: scaine@atersys.com				
	High quality sequence stop: 550.				
FEATURES	Location/Qualifiers				
source	1..803				
	/organism="Homo sapiens"				
	/db_xref="taxon:9606"				
	/clone_lib="Athersys RAGE Library"				
	/cell_line="HT1080"				
	/note="See 'Creation of Genome-wide Protein Expression Libraries using Random Activation of Gene Expression', Nature Biotechnology, in press. Note that even though the cell type indicated is HT1080, since a random activation method was used, these sequence tags are not necessarily expressed in HT1080 under normal circumstances."				



Seq primer: -40UP from GIBCO  
High quality sequence stop: 411.  
Location/Qualifiers  
1. 429

/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:2308895"  
/clone.lib="NCI.CGAP.GC6"  
/tissue.type="pooled germ cell tumors"  
/lab\_host="DH10B"

/note="Vector: p773D-Pac (Pharmacia) with a modified polylinker. Site 1: Not I; Site 2: Eco RI; Plasmid DNA from the normalized library NCI.CGAP.GC4 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clones 1257096-1258631, 1469064-1470983, and 1475592-1476743). Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 127 a 100 c 97 g 104 t 1 others

ORIGIN

Query Match 90.3%; Score 336; DB 9; Length 429;

Best Local Similarity 100.0%; Pred. No. 3.2e-90; Mismatches 0; Indels 0; Gaps 0;

Matches 336; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 37 GAGCTCTCCGAGAGATTTTATTACTTATATGACAAATGGCGCCAGAACACAGCTG 96

Db 12 GAGCTCTCCGAGAGATTTTATTACTTATATGACAAATGGCGCCAGAACACAGCT 71

QY 97 GAGCAAGAGGCGCTCCAGGCAAGTTGATGCTGAGAACTTCTATGTCATCTGTAC 156

Db 72 GAGCAAGAGGCGCTCCAGGCAAGTTGATGCTGAGAACTTCTATGTCATCTGTAC 131

QY 157 CTGATGTCATGATTTGGAATGTCCTTTCATATGTCGTGACCTCTGTGAGACATG 216

Db 132 CTGATGTCATGATTTGGAATGTCCTTTCATATGTCGTGACCTCTGTGAGACATG 191

QY 217 AATTCAGAGAGCGGAGACCTCAATGACCCCTACACAGTACATTTGAGAGACTG 276

Db 192 AATTCAGAGAGCGGAGACCTCAATGACCCCTACACAGTACATTTGAGAGACTG 251

QY 277 CAGGAAAGTACAGAGCCCAATCTTGAATCTGAGAAATCGAAGGCCACATCATGAG 336

Db 252 CAGGAAAGTACAGAGCCCAATCTTGAATCTGAGAAATCGAAGGCCACATCATGAG 311

QY 337 AACATTGGTGGCGCTGGTTCAAAATGTCCTCCCTGA 372

Db 312 AACATTGGTGGCGCTGGTTCAAAATGTCCTCCCTGA 347

RESULT 4 AI339609 391 bp mRNA linear EST 29-DEC-1998

LOCUS AI339609 391 bp mRNA linear EST 29-DEC-1998

DEFINITION q942a07.x1 Soares\_NHMPu.S1 Homo sapiens cDNA clone IMAGE:1935156

3' similar to SW:MINK.HUMAN.P15382 ISK SLOW VOLTAGE-GATED POTASSIUM

CHANNEL PROTEIN ; mRNA sequence.

ACCESSION AI339609.1 GI:4076536

VERSION AI339609.1 GI:4076536

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 391) Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE Tumor Gene Index

JOURNAL Contact: Robert Strausberg, Ph.D.

Seq primer: -40UP from GIBCO  
High quality sequence stop: 380.  
Location/Qualifiers  
1. 391

/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1935156"  
/clone.lib="Soares\_NHMPu.S1"  
/tissue.type="Pooled human melanocyte, fetal heart, and pregnant uterus"  
/lab\_host="DH10B"

/note="Organ: mixed (see below); Vector: p773D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2NbM, pregnant uterus 2NbHU, and fetal heart 2NbH19W) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479."

BASE COUNT 119 a 93 c 93 g 86 t

ORIGIN

Query Match 89.2%; Score 331.8; DB 9; Length 391;

Best Local Similarity 99.4%; Pred. No. 5.7e-89; Mismatches 2; Indels 0; Gaps 0;

Matches 333; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 38 ACGTCTCCGAGAGATTTTATTACTTATATGACAAATGGCGCCAGAACACAGCTG 97

Db 3 ACGTCTCCGAGAGATTTTATTACTTATATGAGGGAATGGCGCCAGAACACAGCTG 62

QY 98 AGCAAGAGGCGCTCCAGGCAAGTTGATGCTGAGAACTTCTATGTCATCTGTAC 157

Db 63 AGCAAGAGGCGCTCCAGGCAAGTTGATGCTGAGAACTTCTATGTCATCTGTAC 122

QY 158 TCATGTCATGATTTGGAATGTCCTTTCATATGTCGTGACCTCTGTGAGACATG 217

Db 123 TCATGTCATGATTTGGAATGTCCTTTCATATGTCGTGACCTCTGTGAGACATG 182

QY 218 AATTCAGAGAGCGGAGACCTCAATGACCCCTACACAGTACATTTGAGAGACTG 277

Db 183 AATTCAGAGAGCGGAGACCTCAATGACCCCTACACAGTACATTTGAGAGACTG 242

QY 278 AGGAAAGTACAGAGCCCAATCTTGAATCTGAGAAATCGAAGGCCACATCATGAG 337

Db 243 AGGAAAGTACAGAGCCCAATCTTGAATCTGAGAAATCGAAGGCCACATCATGAG 302

QY 338 ACATTGGTGGCGCTGGTTCAAAATGTCCTCCCTGA 372

Db 303 ACATTGGTGGCGCTGGTTCAAAATGTCCTCCCTGA 337

RESULT 5 AI246239 372 bp mRNA linear EST 28-JAN-1999

LOCUS AI246239 372 bp mRNA linear EST 28-JAN-1999

DEFINITION q129g04.x1 Soares\_NHMPu.S1 Homo sapiens cDNA clone IMAGE:1857942

3' similar to SW:MINK.HUMAN.P15382 ISK SLOW VOLTAGE-GATED POTASSIUM

CHANNEL PROTEIN ; mRNA sequence.

ACCESSION AI246239

VERSION AI246239

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 372) Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE Tumor Gene Index

JOURNAL Contact: Robert Strausberg, Ph.D.

COMMENT Unpublished (1997)



	CDS		/straln="C57BL/6J" /db_xref="MGJ:1901204" /db_xref="taxon:10090" /clone="2200002116" /sex="male" /tissue_type="stomach" /clone_lib="RIKEN full-length enriched mouse cDNA library" /dev_stage="adult" 149..520 .note="data source:SPFR, source key:O9PWVW0, evidence:ISS homolog to MINIMOM POTASSIUM ION CHANNEL-RELATED PEPTIDE 1 (MIRP1) (MINK-RELATED PEPTIDE 1) putative"
	polyA_signal		/codon_start=-1 /protein_id="BAB25781.1" /note="GI:12842914" /translation="MATLANCTQLEDPAFKIFITYMDSWRRNTAEAOLOARVDAAE NFYVIIIVLMVMIGSEFIYVAIIIVSVKSKRHRSDPYHQYIVEDMQEKYKQILIH LEDSKATIHENMKATGPTTVP" 1673..1678 .note="putative"
	polyA_site		1691 .note="putative"
	BASE COUNT	482 a    365 c    410 g    434 t	
	ORIGIN		
	Query Match	72.5%; Score 269.6; DB 11; Length 1691;	
	Best local Similarity	82.8%; Pred. No. 5.4e-70;	
	Matches 308; Conservative 0; Mismatches 64; Indels 0; Gaps 0;		
Qy	1 ATGTCTACTTTATTCATTTCACACAGCGGTGGAGAAGCGTCCGAAGCATTTTATT 60		
Db	149 ATGGCCACATTAGGCCAATTGACCAGACACTGGAGATGCCTTCMAAAGCATTTTATT 208		
Oy	61 ACTTATATGGAACAATTTGGCGCAAGAACCAACAGCTGAGCAAGGCCCTCCAAGCCAAA 120		
Db	209 ACTTATATGGAACAGCTGGAGAGGAACACGACGCCGAGAGCACCTCAGGCCACGA 268		
Oy	121 GTTGATGCTGGAACCTTCTACTATGTCACTGTAACCTCATGSGTAGATTGGAATGTTTC 180		
Db	269 GTGATGCGCGAAGACTTCTACTAGCTCANCTGTACTCATGSGTAGATGCGGATGTTTC 328		
Oy	181 TCTTTCATCATCGTGGCCATCTGCTGTGACACTGTGAAATCCAGACAGCGGACACTCC 240		
Db	329 TCGTTCATCGGTGGCGCATCTGCTGTAGCACGGGTGAAGTGCMAACGGCGGAGACACTCC 388		
Oy	241 AATGACCCCTGCACCACTACATGTGTAGAGAGCTGGAGAAAAGTAGAAGCCCAATTC 300		
Db	389 CAGGACCCCTGCACCACTACATCTGTGAAGTTTGGCAGAAAAATACAAAAGTCAGATTC 448		
Oy	301 TTGATCTAGAAGAAATCCAAAGCCACCATCATGAGAACATTGTGCGGCTGCTTCAA 360		
Db	449 CTGATCTGGAAGAACTCCAAAGCCACCATCATGAGAACATGSGGGGCGACGGGGTTCACA 508		
Oy	361 ATGTCCCCCTGA 372		
Db	509 GTGTCAACCTGA 520		
RESULT 7			
LOCUS	D85797	470 bp	mRNA linear EST O9-SEP-1996
DEFINITION	D85797 Rat 21 day old female ovary mRNA PM5G 3h Rattus norvegicus		
ACCESSION	D85797		
VERSION	D85797.1		
KEYWORDS	EST.		
SOURCE	Norway rat.		
ORGANISM	Rattus norvegicus		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus. 1 (bases 1 to 470)		

AUTHORS	Miyamoto,K., Mizutani,T., Nunata,Y. and Okada,T.
TITLE	Rat 21 day old female ovary mRNA
JOURNAL	Unpublished (1996)
COMMENT	Contact: Miyamoto,K. . Institute for Molecular and Cellular Reg Gunma Univ., Biosignal Research Center Shouna machi 3-39-15, Maebashi, Gunma 371, Japan Email: umizuesb.gunma-u.ac.jp 5'-59.
FEATURES	
source	Location/Qualifiers 1..470 /organism="Rattus norvegicus" /strain="qistar" /db_xref="taxon:10116" /clone_lib="Rat 21 day old female ovary mRNA PMSG 3h" /sex="female" /tissue_type="ovary" /dev_stage="21 day old"
BASE COUNT	116 a 127 c 128 g 95 t 4 others
ORIGIN	
Query Match	70.1%; Score 260.8; DB 10; Length 470;
Best Local Similarity	80.9%; Pred.No.1.4e-67;
Matches 301; Conservative	0; Mismatches 71; Indels 0; Gaps 0;
OY	1 ATGCTACTTATTCACAAATTTCACACAGACGTGGAGAAGCGTTCCGAAGCATTTTATT 60   Db 37 ATGACCACCTTAGGCCAAGCTTGACGCAGACCCCTGGAGAGATGCCCTCAAAAAGGTTTTTCATT 96  OY 61 ACTTATTAAGACAATAITGGCGCCAGAACCAACAGCTGAGCAMAAGAGCCCCCTCCAACCCCAA 120   Db 97 AATTATTAAGACAGCTGAGGAGGAGCACCAACAGSCGAACCAACAGCCCCTCCAGGCCAGAA 156  OY 121 GTTGAATGCGAGAACTTCTACTATGTCAATCCCTGTAAGCTCATGATGATGATGGATGTTTC 180   Db 157 GTGGATGCCGGAACATTCTACTAGCTCATCCGTGTAAGCTCATGTGATGATGCGCATTTTC 216  OY 181 TCTTTCAATCANTGTGGCCATCTGTGTGAGCACTGTGAANATCCAGAAGCGGAACTTCC 240   Db 217 GCCTTCATCTGTGTGGCCATCTGTGTGAGCAAGGTGAAGTCGAACCGCGGAGCAGCATTC 276  OY 241 ATTGACCCCTACCCACAGTACATTTGTAGAGAGCTGGCAGAAAAATPACAAGGCCAATTC 300   Db 277 CAGGACCCGTACCACAGTNCMTGTGGAGATTGGCAGCAAGATATAGAGGTAGATC 336  OY 301 TTGAATCTAAGAAMTCGAAAGCCACCAATCCATGTAGAAACATGTGTGCGGCTGGTTCAAA 360   Db 337 TTGCAATCTGGAAGACTCCAAAGGCCATCATCATGATTAACCTGGGGGCGACGGGGTTCCACA 396  OY 361 ATGTCCOCCTGA 372                Db 397 NTGTCAACCTGA 408
RESULT 8	
Bg261965	
LOCUS	BG261965 1003 bp mRNA linear EST J3-FEB-2001
DEFINITION	602373784.T NIH_MGC_94 Mus musculus CDNA clone IMAGE:4481325 5',
mRNA sequence.	
Accession	BG261965
VERSION	BG261965.1 GI:12771781
KEYWORDS	EST.
SOURCE	house mouse.
ORGANISM	Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	
NIH-MGC http://mgi.nci.nih.gov/ National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999) Contact: Robert Stransberg, Ph.D. Email: cgabs+remail.nih.gov Tissue Procurement: The Cepko Laboratory	

CDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://www.llnl.gov/imagetdb>

Plate: L1AM10316 row: 9 column: 22  
High quality sequence stop: 535.

FEATURES	Location/Qualifiers
source	1. .1003

BASE COUNT	249 a	218 c	302 g	234 t
ORIGIN				

Query Match	65.9%; Score 245; DB 10; Length 1003;
-------------	---------------------------------------

Matches 307; Conservative 0; Mismatches 65; Indels 3; Gaps 2;

1 ATGTCTACTTTATCCAAITTCACACAGACGCTGGAGAAGACGTTCCGAAGATTTTATT 60

91 A TGGCACAATTAGCCAAATTGACCCAGACACTGGAGGATGCCCTTCAAAAAGATTTTATT 150

01 ACCTTATGGACAACTGGCCCGAGAACACACACACAGCCTGAGCAAGAGGCCCTCCAAAGCCAA 120

101 ACCTATGATACGCTGGAGGAGGACGACCTCCAGGCCAGA 210

[illegible][illegible]

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[illegible]

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**301**

[illegible]

358 AAAATGTCGCCCTGA 372

451 ACAGTGTCAACCCTGA 465

RESULT 9			
LOCUS	746 bp	mRNA	linear
DEFINITION	ATHERSYS RAGE Library Homo sapiens cDNA, mRNA sequence.		
ACCESSION	U02119		
VERSION	U02119.1		
KEYWORDS			
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ORIGIN	99		
ORIGIN	100		

ORGANISM	Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.	

**TITLE** ,J., Denzly,J. and Ducrar,M.  
Creation of genome-wide protein expression libraries using random  
activation of gene expression  
**JOURNAL** Nat. Biotechnol. 19 (5), 440-445 (2001)  
**ADDRESS**

COMMENT	CONTACT: SCOTT J. CALIN	ATTORNEY: TONY

3201 Carnegie Ave, Cleveland, OH 44115, USA  
Tel: 216 431 9900

Email: [scain@athersys.com](mailto:scain@athersys.com)  
High quality sequence atop: 547

FEATURES	Location/Qualifiers
SOURCE	1. .746

BASE COUNT	241 a	138 c	123 g	243 t	1 others
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Best Local Similarity 93.2%; Pred. No. 1.5e-42;

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Db 586 ACTTATATGACAATTGGCCGACACACACCACTGAGCAGGAGGCCCTCC-AGCCAAA 644

121 GTTGATGCTGAGAACTTCTACTATGTGCATCCTGTACCTCATGGTGATGATTGGAATGTC 180

Db 645 GTTGATGCTGAGAACTTCTACTATGTTCATCCCGACCAATGNGATGATTGGAATGTTTC 704

181 TCTTTCAT-CATCGTGGCCATCCTGGTGAGCACTGTGAATC 221  
QY

	RESULT 10	
BG938225		
LOCUS		
BG938225	351 bp	mRNA linear EST JJ-JUN-2001

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.

**COMMENT**  
Contact: Dr. Stephen Moore  
. Beef Genomics Laboratory  
Dept of AFNS, University of Alberta

chromosome 21q, section 63/105) in main database at high score of 212.0 and E-value of 9e-53  
 PCR Primers  
 FORWARD: M13 Forward  
 BACKWARD: M13 Reverse  
 Seq primer: T3 primer  
 High quality sequence stop: 351  
 POLYA-No.

FEATURES  
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/organism="Bos taurus"  
 /db\_xref="taxon:9913"  
 /clone\_lib="Bovine Abomasum cDNA Library"  
 /sex="Two males and one female mixed"  
 /tissue\_type="Gastrointestinal tissue (GIT)"  
 /cell\_type="Epithelial"  
 /dev\_stage="young adult"  
 /lab\_host="XLI-BlueMRF"-strain"  
 /note="Organ: Abomasum; Vector: Uni-22APYR; Site\_1: EcoR I; Site\_2: Xho I"

BASE COUNT 104 a 89 c 76 g 82 t  
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Query Match 45.2%; Score 168.2; DB 10; Length 351;  
 Best Local Similarity 87.1%; Pred. No. 8.4e-40;  
 Matches 196; Conservative 0; Mismatches 28; Indels 1; Gaps 1;

OY 1 ATGTACTCTTATCCAAATTCACAGACGCTGGAAAGCGTTCCGAAGATTTTATT 60  
 DB 127 ATGCCAAGCTATCCAAATTCACAGACGCTGGAAATATGCTTCAAAAATTTTATC 186  
 OY 61 ACTTATATGACAAATTCGCGCAGACAAACAGCTGAGCAAGAGCCCTCCAGCCAA 120  
 DB 187 ACTTATATGACAAATTCGCGCAGACAAACAGCTGAGCAAGAGCCCTCCAGCCAA 246  
 OY 121 GTTATGCTGAGAACTTCTACTATGTCACTCTGACCTCAGTGATGTAATGTTTC 180  
 DB 247 GTTATGCTGAGAACTTCTACTATGTCACTCTGACCTCAGTGATGTAATGTTTC 306  
 OY 181 TCTTTCATCCTGCGCAGCTGAGCAAGCTGTAATCCAA 224  
 DB 307 TCTTTCATCCTGCGCAGCTGAGCAAGCTGTAATCCAA 351

RESULT 11 188 bp mRNA linear EST 28-OCT-1997  
 AA633404  
 LOCUS npe9h11.s1 NCI-CGAP Br2 Homo sapiens cDNA clone IMAGE:1131621 3'  
 DEFINITION similar to SW:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.

ACCESSION AA633404  
 VERSION AA633404.1  
 KEYWORDS GI:2555264  
 SOURCE EST.  
 ORGANISM human.

REFERENCE Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 TITLE Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.  
 Emmert-Buck, M.D., Ph.D.  
 cDNA Library Preparation: M. Bento Soares, Ph.D.  
 DNA Sequencing by: Greg Lennon, Ph.D.  
 Clone distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 www-bio.llnl.gov/bdrip/image/image.html  
 Insert Length: 785 Std Error: 0.00

Seq primer: -40m13 fwd. ET from Amersham  
 High quality sequence stop: 167.

FEATURES  
 source Location/Qualifiers  
 1.188

/organism="Homo sapiens"  
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 /clone\_lib="IMAGE:1131621"  
 /clone\_lib="NCI-CGAP Br2"  
 /sex="female, pooled"  
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 /lab\_host="DH10B"  
 /note="Vector: p773D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from pooled bulk breast tumor tissue, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p773 vector. This library is the normalized version of NCI CGAP Br1.1. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 66 a 45 c 45 g 32 t  
 ORIGIN

Query Match 43.0%; Score 160; DB 9; Length 188;  
 Best Local Similarity 100.0%; Pred. No. 1.9e-37;  
 Matches 160; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 213 TGTGAATCCAGAGAGCGGACACCTCAATGACCCCTACCAAGATGTTAGAGA 272  
 DB 1 TGTGAATCCAGAGAGCGGACACCTCAATGACCCCTACCAAGATGTTAGAGA 60  
 OY 273 CTGGCAGAAAGATGACAGACCAATCTGAAAGATGGAAGGCACATCCCA 332  
 DB 61 CTGGCAGAAAGATGACAGACCAATCTGAAAGATGGAAGGCACATCCCA 120  
 OY 333 TGAGAACATGCTGCTGGCTGCTCAAAATGTCCCTGCA 372  
 DB 121 TGAGAACATGCTGCTGGCTGCTCAAAATGTCCCTGCA 160

RESULT 12 121 bp mRNA linear EST 07-JUL-1998  
 AA935321  
 LOCUS o071g09.s1 NCI-CGAP GC4 Homo sapiens cDNA clone IMAGE:1571680 3'  
 DEFINITION similar to SW:MINK\_HUMAN P15382 ISK SLOW VOLTAGE-GATED POTASSIUM CHANNEL PROTEIN ; mRNA sequence.

ACCESSION AA935321  
 VERSION AA935321.1  
 KEYWORDS GI:3092478  
 SOURCE EST.  
 ORGANISM human.

REFERENCE Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 TITLE Tumor Gene Index  
 JOURNAL Unpublished (1997)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgaps-remail.nih.gov  
 Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael  
 Emmert-Buck, M.D., Ph.D.  
 cDNA Library Preparation: M. Bento Soares, Ph.D.  
 DNA Sequencing by: Greg Lennon, Ph.D.  
 Clone distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 www-bio.llnl.gov/bdrip/image/image.html  
 Trace considered overall poor quality  
 Insert Length: 763 Std Error: 0.00  
 Seq primer: -40m13 fwd. ET from Amersham  
 High quality sequence stop: 1.

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source  
Location/Qualifiers  
1. 121  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_id="IMAGE:1571680"  
/issue\_type="pooled germ cell tumors"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from 3 pooled germ cell tumors, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT  
ORIGIN  
33 a 31 c 28 g 29 t

Query Match  
Best Local Similarity 99.2%; Pred. No. 2,5e-25;  
Matches 120; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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OY 213 TGTGAATTCAGAGGAGGAGACACCTCATCATGACCCCTACACAGATGATTTGAGAGA 272  
|||||  
Db 61 TGTGAATTCAGAGGAGGAGACACCTCATCATGACCCCTACACAGATGATTTGAGAGA 120

OY 273 C 273  
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Db 121 C 121

RESULT 13  
AM869303 311 bp mRNA linear EST 22-MAY-2000  
LOCUS  
DEFINITION  
MR3-SN0067-240400-006-f11 SN0067 Homo sapiens cDNA, mRNA sequence.  
AM869303  
VERSION  
AM869303.1 GI:8003356  
KEYWORDS  
EST.  
SOURCE  
human.  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE  
AUTHORS  
1 (bases 1 to 311)  
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bala,G.S., Simpson,D.H., Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.  
Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
20202663  
JOURNAL  
MEDLINE  
COMMENT  
Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=612-MR3-SN0067-240400-006-f11&ts=2000-04-24&td=1)  
Seq primer: puc 18 forward  
High quality sequence start: 11  
High quality sequence stop: 74.

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source  
Location/Qualifiers  
1. 311  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone\_id="SN0067"  
/dev\_stage="Adult"  
/note="Organ: stomach:normal; Vector: puc18; Site-1: SmaI; Site-2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

BASE COUNT  
ORIGIN  
60 a 80 c 76 g 95 t

Query Match  
Best Local Similarity 90.7%; Pred. No. 5.1e-19;  
Matches 117; Conservative 0; Mismatches 10; Indels 2; Gaps 1;

OY 1 ATGCTACTTTATTCATTTACACAGACGCTGGAAGAGCTCTCCGAGATTTTAT 60  
|||||  
Db 182 ATGCTACTTTATTCATTTACACAGACGCTGGAAGAGCTCTCCGAGATTTTATC 123

OY 61 ACTTATATGACAAATTGG--CGCCAGACACACACACCTGAGAGAAGCCCTCAAGCCA 118  
|||||  
Db 122 ACTCATATGACAAATGGCGCGCGAGACACGACCTGAGCGAGAGCCCTCAAGCCT 63

OY 119 AAGTTGATG 127  
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Db 62 AAGATGATG 54

RESULT 14  
BB564873 314 bp mRNA linear EST 29-NOV-2000  
LOCUS  
DEFINITION  
BB564873 RIKEN full-length enriched, adult male stomach Mus musculus cDNA clone 220002116 5', mRNA sequence.  
BB564873  
VERSION  
BB564873.1 GI:11455765  
KEYWORDS  
EST.  
SOURCE  
house mouse.  
ORGANISM  
Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE  
AUTHORS  
1 (bases 1 to 314)  
Aizawa,K., Akahira,S., Akimura,T., Arai,A., Arakawa,T., Carolincl,P., Hanagaki,T., Hayatsu,N., Hirooka,T., Hirozane,T., Hodyama,Y., Imotani,K., Ishii,Y., Itoh,M., Izawa,M., Kawai,J., Kojima,Y., Konno,H., Kusakabe,M., Matsuyama,T., Miyazaki,A., Nakamura,M., Nishi,K., Nomura,K., Nunakazi,K., Okazaki,Y., Okido,T., Owa,C., Sakai,C., Sakai,K., Sasaki,D., Sato,K., Shibata,K., Shibata,Y., Shinagawa,A., Shiraki,T., Sogabe,Y., Iizuki,H., Tategawa,A., Takahashi,F., Tanaka,T., Toyota,T., Watabiki,A., Yamamura,T., Yasunishi,A., Yoshida,K., Yoshiki,A., Muramatsu,M. and Hayashizaki,Y.  
RIKEN Mouse ESTs (Aizawa,K. et al. 2000)  
Unpublished (2000)  
Contact: Yoshihide Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suenho-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216  
Email: genome-res@sc.riken.go.jp/  
URL: http://genome.gsc.riken.go.jp/  
Carlincl,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.  
Thermolabile and thermocatalytic activation of thermolabile enzymes by thermolabile and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998)  
Itoh,M., Kitsumai,T., Akiyama,J., Shibata,K., Izawa,M., Kawai,J., Tomaru,Y., Carolincl,P., Shibata,Y., Ozawa,Y., Muramatsu,M., Okazaki





Tue May 27 09:59:42 2003

us-09-550-163-1\_copy\_74\_445.rst

Page 10

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Job time : 1610 secs

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GenCore version 5.1.4.p5\_4578  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 20:14:00 ; Search time 21.4496 Seconds

(without alignments)  
5275.799 Million cell updates/sec

Title: US-09-550-163-1\_COPY\_74\_442

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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Listing first 45 summaries

Database : Issued\_Patents\_NA:\*  
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4: /cgn2\_6/ptodata/1/lna/6B.COMB.seq:\*  
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Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	53.2	14.4	398	1	US-08-118-101A-5
2	53.2	14.4	436	4	US-09-679-185-1
3	53.2	14.4	1703	3	US-09-135-021-77
4	53.2	14.4	1703	4	US-09-135-020-3
5	53.2	14.4	1703	4	US-09-135-010A-3
6	53.2	14.4	1703	4	US-09-444-871-3
7	53.2	14.4	1703	4	US-09-597-733-3
8	53.2	14.4	1703	4	US-09-444-295-3
9	53.2	14.4	1703	4	US-09-597-733-3
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11	36.6	9.9	2652	1	US-08-318-831-1
12	36	9.8	1380	1	US-08-110-286A-1
13	36	9.8	1495	4	US-08-482-746-1
14	36	9.8	1582	4	US-08-482-746-14
15	33.6	9.1	645	4	US-09-069-896-2
16	33.6	9.1	645	4	US-09-471-468-2
17	33	8.9	606	4	US-09-328-111-133
18	33	8.9	2812	4	US-09-307-143-3
19	32.4	8.8	837	4	US-08-998-416-303
20	32.4	8.8	8659	4	US-09-221-017B-823
21	30.6	8.3	6822	4	US-09-426-998-3
22	30.6	8.3	7741	4	US-09-961-527-83
23	30.2	8.2	28473	4	US-09-215-694-18
24	30.2	8.2	33000	4	US-08-724-394A-18
25	29.8	8.1	2266	2	US-08-724-394A-20
26	29.8	8.1	246240	2	US-08-724-394A-21
27	29.8	8.1	246240	2	US-08-724-394A-21

28	29.8	8.1	246240	2	US-08-724-394A-22	Sequence 22, Appl
29	29.6	8.0	462	3	US-08-863-813A-33	Sequence 33, Appl
30	29.6	8.0	3273	6	US-09-630-1	Patent No. 551630
31	29.6	8.0	6002	4	US-09-345-882-4	Sequence 4, Appl1
32	29.6	8.0	7486	3	US-08-475-886-5	Sequence 5, Appl1
33	29.6	8.0	7486	4	US-08-397-232-3	Sequence 3, Appl1
34	29.6	8.0	7486	4	US-09-653-499-5	Sequence 5, Appl1
35	29.6	8.0	7486	3	US-08-475-886-3	Sequence 3, Appl1
36	29.6	8.0	7488	4	US-09-653-499-3	Sequence 3, Appl1
37	29.6	8.0	7493	4	US-08-475-886-1	Sequence 1, Appl1
38	29.6	8.0	7493	4	US-08-397-232-1	Sequence 1, Appl1
39	29.6	8.0	7493	4	US-09-171-387-1	Sequence 1, Appl1
40	29.6	8.0	7493	4	US-09-653-499-1	Sequence 1, Appl1
41	29.4	8.0	1666	4	US-09-221-017B-933	Sequence 933, App
42	29.4	8.0	3947	4	US-08-975-762-47	Sequence 47, Appl
43	29.4	8.0	3947	4	US-09-235-028-47	Sequence 47, Appl
44	29.4	8.0	3947	4	US-09-106-582-47	Sequence 47, Appl
45	29.2	7.9	352	4	US-09-605-785-421	Sequence 421, App

#### ALIGNMENTS

RESULT 1  
US-08-118-101A-5  
; Sequence 5, Application US/08118101A  
; Patent No. 5620892  
; GENERAL INFORMATION:  
; APPLICANT: Kuitz, Stephen E.  
; APPLICANT: Knickerbocker, Aron M.  
; APPLICANT: McCullough, John R.  
; TITLE OF INVENTION: A STRAIN OF SACCAROMYCES CEREVISIAE  
; TITLE OF INVENTION: EXPRESSING THE GENE ENCODING POTASSIUM TRANSPORTER MINK  
; NUMBER OF SEQUENCES: 16  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Burton Rodney  
; STREET: P.O. Box 4000  
; CITY: Princeton  
; STATE: New Jersey  
; COUNTRY: U.S.A.  
; ZIP: 08543-4000  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/118,101A  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Gaul, Timothy J.  
; REGISTRATION NUMBER: 33,111  
; REFERENCE/DOCKET NUMBER: DC27  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (609) 252-5901  
; TELEFAX: (609) 252-4526  
; INFORMATION FOR SEQ ID NO: 5:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 398 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: cDNA  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 1..398  
; US-08-118-101A-5  
Query Match 14.4%, Score 53.2, DB 1; Length 398;  
Best Local Similarity 63.6%, Pred. No. 1,7e-08;  
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

QY 210 CACTGTGAATCCAAAGACGGGAAACACTCCATGACCCCTACACCAAGTAAATTG---T 266  
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 QY 150 CCGTACCTCATATGGTATGATGTGAATGTTCTCTTTATCATCTGTGGCAATCCGTGGAG 209  
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 QY 267 AGAGGACTGGGAGGAAAAAGTACAGAGACCAATC 300  
 Db 261 CGATGCTCGGCMAAGAGAGCAAGCAAGGCGCTATGTC 294

RESULT 2  
NE-09-67

Sequence 1, Application US/09679185  
Patent No. 6458542  
GENERAL INFORMATION:  
APPLICANT: George Jr., Alfred L.  
APPLICANT: Roden, Dan M  
TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO  
TITLE OF INVENTION: DRUG-INDUCED CARDIAC ARRHYTHIA  
FILE REFERENCE: Attorney Docket No. 6458542 1242-33-2  
CURRENT APPLICATION NUMBER: US/09/679,185  
CURRENT FILING DATE: 2000-10-04  
PRIOR APPLICATION NUMBER: 60/158,696  
PRIOR FILING DATE: 1999-10-08  
NUMBER OF SEQ ID NOS: 11  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 1  
LENGTH: 436  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (29)..(418)  
PUBLICATION INFORMATION:  
JOURNAL: Biochem. Biophys. Res. Commun.  
VOLUME: 161  
PAGES: 176-181  
DATE: May-1989  
DATABASE ACCESSION NUMBER: GenBank M26685  
DATABASE ENTRY DATE: 1994-03-30  
US-09-679-185-1

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Dy	220 CTACATTCGCCGCTCCAGAAAGACTGAGCACATCTAGACAGCACCATTCAACCTTACATCGAGTC				279
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Dy	280 CGATGCTTGGCAAAGAAAGAACGACCAAGGCTTAATGTC				313

### RESULT 3

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US-09-135-021-77
; Sequence 77, Application US/09135021A
; Patent No. 6150104
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; GENERAL INFORMATION:
; APPLICANT: SPlawski, Igor
; APPLICANT: Keating, Mark T.
; TITLE OF INVENTION: A HOMOLOGOUS MUTATION IN KVLQ1 WHICH CAUSES JERVELL
; TITLE OF INVENTION: AND LANGE-NIELSEN SYNDROME
; FILE REFERENCE: 2323-128
; CURRENT APPLICATION NUMBER: US/09/135.021A

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: CURRENT FILING DATE: 1998-08-17
: EARLIER APPLICATION NUMBER: 08/874,655
: EARLIER FILING DATE: 1997-06-13
: EARLIER APPLICATION NUMBER: 60/094,477
: EARLIER FILING DATE: 1998-07-29
: NUMBER OF SEQ ID NOS: 80
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 77
: LENGTH: 1703
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: CDS
: LOCATION: (193)..(579)
US-09-135-021-77

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Query Match  
Post Total

Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

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Db	324	CCTGACGCTCTCATGGTACTGGGATTTCTTGCGGCTTTTCAACCTGGGCATCATGCTGAG	383
QY	210	CACGTGAAATCCAAAGAGACGGGAAACACTCCAAAGACCCCTAACCAACAGTAAATG---T	266
Db	384	CTACATCCGCTCCAAAGACCTGGGACCTCTCAAGACCCATCTCAACGCTTACGAGATC	443
QY	267	AGAGGACTGCGCAGGAAAGTACAAAGAGCCCAATC	300
Db	444	CGATGCTCTGGCCAGAGAAAGGACCAAGGCGCTTATGTC	477

## RESULT 4

Sequence 3 Application US/09135020  
 Patient No. 6274332  
 GENERAL INFORMATION:  
 APPLICANT: Keating, Mark T.  
 APPLICANT: Sanguinetti, Michael C.  
 APPLICANT: Splawski, Igor  
 TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MINK WHICH  
 TITLE OF INVENTION: CAUSE ARRYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING  
 TITLE OF INVENTION: KCNE1 AS AN LQT GENE  
 FILE REFERENCE: 2323-131  
 CURRENT APPLICATION NUMBER: US/09/135,020  
 CURRENT FILING DATE: 1998-08-17  
 EARLIER APPLICATION NUMBER: 08/921,068  
 EARLIER FILING DATE: 1997-08-29  
 EARLIER APPLICATION NUMBER: 08/739,383  
 EARLIER FILING DATE: 1996-10-23  
 EARLIER APPLICATION NUMBER: 60/019,014  
 EARLIER FILING DATE: 1995-12-22  
 EARLIER APPLICATION NUMBER: 60/094,477  
 EARLIER FILING DATE: 1998-07-29  
 NUMBER OF SEQ ID NOS: 114  
 SOFTWARE: Patentln Ver. 2.0

TYPE

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:   FEATURE:
:   NAME/KEY: CDS
:   LOCATION: (193)..(579)
US-09-135-020-3

Query Match          14.4%   Score 53.2;   DB 4;   Length 1703;
Best Local Similarity 63.6%   Pred. No. 3.9e-08;
Matches 98; Conservative 0; Mismatches 53; Indels 3; Gaps 1

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OY     150 CCGTACCCATGGGATGATTGGAAGTTCTTCCTTCAATCATGCnGCCAACCCTGGTAG   209  
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Db     324 CCTCTACGCCCTCATTGTACTGGGATCTTGCGCTTCTTCACCCCTGGGCATCATGCTGAG   383

; TITLE OF INVENTION: CHOSE ANNIHILATA SUSCEPTIBILITY INHEKBI ESTABLISHING  
; TITLE OF INVENTION: KCNE1 AS AN IQT GENE

ORGANISM: Homo sapiens

```

;
;
; TYPE: DNA
; ORGANISM: Homo sapiens

```

```

: RESULT 7
: US-09-597-735-3
: Sequence 3, Application US/09597735
: Patent No. 6420124
: GENERAL INFORMATION:
: APPLICANT: Keating, Mark T.
: APPLICANT: Sanguinetti, Michael C.
: APPLICANT: Curran, Mark E.
: APPLICANT: Landes, Gregory M.
: APPLICANT: Connors, Timothy D.
: APPLICANT: Burn, Timothy C.
: APPLICANT: Splawski, Igor
: TITLE OF INVENTION: KVLQTL - A LONG QT SYNDROME GENE
: FILE REFERENCE: 2323-113
: CURRENT APPLICATION NUMBER: US/09/597,735
: CURRENT FILING DATE: 2000-06-19
: EARLIER APPLICATION NUMBER: 09/135,010
: EARLIER FILING DATE: 1998-08-17
: EARLIER APPLICATION NUMBER: 60/094,477
: EARLIER FILING DATE: 1998-07-29
: EARLIER APPLICATION NUMBER: 08/921,068
: EARLIER FILING DATE: 1997-08-29
: EARLIER APPLICATION NUMBER: 08/739,383
: EARLIER FILING DATE: 1996-10-29
: EARLIER APPLICATION NUMBER: 60/019,014
: EARLIER FILING DATE: 1995-12-22
: NUMBER OF SEQ ID NOS: 116
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 3
: LENGTH: 1703
: TYPE: DNA
: ORGANISM: Homo sapiens

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; FEATURE:
; NAME/KEY: CDS
; LOCATION: (193)..(579)
US-09-597-735-3

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Query Match	14.48;	Score 53.2;	DB 4;	length 1703;
Best Local Similarity	63.68;	Pred. No. 3.9e-08;		
Matches 98; Conservative	0;	Mismatches 53;	Indels 3;	Gaps 1;

QY	150	CCTTACCTCATGATGATGAAATGTTCTTTTATCATACGTGGCACCCTGGGGAG	209
Db	324	CCTTACGTCCTCATGAGTACGGGATTTCTTGCGGTTTTCACCTGGGCATCATGCTGAG	383
QY	210	CACGTGAAATCCCAAGACGGGAAACACTCCAAATGACCCTTACACCAAGTACATG---T	266
Db	384	CTACATCCGCTCCAAAGACGTGGAGCATCTGACAGGACCATTCACATGCTTACGAGTC	443
QY	267	AGAGGACTGCGCAGAAAAGTACAAAGACCCAAATC	300
Db	444	CGATGCTGGCCAGAGAGGACCAAGGGCTTATGTC	477

RESULT 8  
US-09-444-295-3  
: Sequence 3, Application US/09444295

? GENERAL INFORMATION:  
 ? APPLICANT: Keating, Mark T.  
 ? APPLICANT: Sanguinetti, Michael C.  
 ? APPLICANT: Splawski, Igor  
 ? TITLE OF INVENTION: MUTATIONS IN THE KCNE1 GENE ENCODING HUMAN MINK WHICH  
 ? TITLE OF INVENTION: CAUSE ARRHYTHMIA SUSCEPTIBILITY THEREBY ESTABLISHING  
 ? TITLE OF INVENTION: KCNE1 AS AN IOT GENE

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; ORGANISM: Homo sapiens
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; FEATURE:
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; NAME/KEY: CDS
; LOCATION: (193)..(579)
;
US-09-444-295-3

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Query Match	Similarity	Score	DB	Length
Best Local	63.6%	Pred. No. 3.9e-08		1103
Matches	98	Conservative	0	Mismatches 53; Indels 3; Gaps 1
QY	150	CCTGTACCTCATGTGATGATTTGTAATGTCTCTTTTCATCTGTCGGCCATCCGTGGAG	209	
Db	324	CCTCATCCGCTCCATCATGTGATGGAATTTCTTGGCTTTTCACTCGGACATCATGCTGAG	383	
QY	210	CACGTGAAATCCAAAGACGGGAACTCCCAATGACCCCTACACAGATGATTTG--T	266	
Db	384	CTATACCTCCGCTCCAAAGACGTGGAGCACTCGAAGACCAATTCATCACTCATTCAGATC	443	
QY	267	AGAGGACTGGCAGAGAAAGTACAAAGGCCAATC	300	
Db	444	CGATGCTGTGGCAAGAGAGACCAAGGCTTATGTC	477	

RESULT 9  
US-09-597-732-3  
; Sequence 3, Application US/09597732

```

1 GENERAL INFORMATION:
2 APPLICANT: Keating, Mark T.
3 APPLICANT: Sanguinetti, Michael C.
4 APPLICANT: Curran, Mark E.
5 APPLICANT: Landes, Gregory M.
6 APPLICANT: Connors, Timothy D.
7 APPLICANT: Burn, Timothy C.
8 APPLICANT: Splawski, Igor
9 TITLE OF INVENTION: KvLQ1 - A LONG QT SYNDROME GENE
10 FILE REFERENCE: 2323-1133
11 CURRENT APPLICATION NUMBER: US/09/597,732
12 CURRENT FILING DATE: 2000-06-19
13 PRIOR APPLICATION NUMBER: 09/135,010
14 PRIOR FILING DATE: 1998-08-17
15 PRIOR APPLICATION NUMBER: 60/094,477
16 PRIOR FILING DATE: 1998-07-29
17 PRIOR APPLICATION NUMBER: 08/921,068
18 PRIOR FILING DATE: 1997-08-29
19 PRIOR APPLICATION NUMBER: 08/739,383
20 PRIOR FILING DATE: 1996-10-29
21 PRIOR APPLICATION NUMBER: 60/019,014
22 PRIOR FILING DATE: 1995-12-22
23 NUMBER OF SEQ ID NOS: 116
24 SOFTWARE: PatentIn Ver. 2.10
25 SEQ ID NO 3

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Query Match	14.48;	Score 53.2;	DB 4;	Length 1703;
Best Local Similarity	63.68;	Pred. No. 3.9e-08;		
Matches 98; Conservative	0;	Mismatches 53;	Indels 3;	Gaps 1

QY	150	CGTGTCCCTCAGTGGATGTGGAAATGTTCTCTTTCATCATCGAGGCCATCGTGGTAG	205
Db	324	CGCTACGTCCTCAGTGGACTGGGATTTCTTGCGGCTTCTTACCCCTGGGGATCATGCTAG	383
QY	210	CAGTGGAAATCCCAAGAACCGGAAACACTCCATGACCCCTACACACAGTACATTG--T	266
Db	384	CTACATCCGCTCCAAAGAGCTGGAGCACTGGAAAGACCCCATTCATCAAGTCTACATGAGTC	443
QY	267	AGAGAGCTGGCAGGAAAGTACAAAGCCCAATTC	300
Db	444	CGATGCTCTGGCAAGAAAGGCAAGGCTTATGTC	477

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1 RESULT 10
2 US-09-679-185-3
3 : Sequence 3, Application US/09679185
4 Patent No. 6458542
5 :
6 : GENERAL INFORMATION:
7 :
8 : APPLICANT: George Jr., Alfred L.
9 :
10 : APPLICANT: Roden, Dan M
11 :
12 : TITLE OF INVENTION: METHOD OF SCREENING FOR SUSCEPTIBILITY TO
13 :
14 : TITLE OF INVENTION: DRUG-INDUCED CARDIAC ARRYTHMIA
15 :
16 : FILE REFERENCE: Attorney Docket No. 6458542 1242-33-2
17 :
18 : CURRENT APPLICATION NUMBER: US/09/679,185
19 :
20 : CURRENT FILING DATE: 2000-10-04
21 :
22 : PRIOR APPLICATION NUMBER: 60/158,656
23 :
24 : PRIOR FILING DATE: 1999-10-08
25 :
26 : NUMBER OF SEQ ID NOS: 11
27 :
28 : SOFTWARE: PatentIn Ver. 2.0
29 :
30 : SEQ ID NO 3
31 :
32 : LENGTH: 436

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: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: CDS
: LOCATION: (29)..(418)
:
US-09-679-185-3

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Query Match	14.08;	Score 51.6;	DB 4;	Length 436;
Best Local Similarity	63.08;	Pred. No. 6.2e-08;		
Matches 97; Conservative	0;	Mismatches 54;	Indels 3;	Gaps 1;

Qy	210	CACGTGAAATCCAAAGAGAGGGGAAACCTCAATGACCCTTACACCAAGATGATGTTG---T	266
Db	220	CTAATCCGGCTCAAGAAAGCTGGAGCACTGACAGACCATTCAAGCTTCATCATGAGTC	275
Qy	267	AGAGGACTGGCAGGAAAAATGACAAAGAGCCCAATC	300
Db	280	CAATGCTGGCAAGAGAAAGGCAAGGGCTTATGTC	313

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US-08-318-831-1
: Sequence 1, Application US/08318831
: Patent No. 5656595
: GENERAL INFORMATION:
: APPLICANT: Schweighoffer, Fabien
: APPLICANT: Tocque, Bruno
: TITLE OF INVENTION: PEPTIDES HAVING A GDP EXCHANGE FACTOR
: TITLE OF INVENTION: ACTIVITY, NUCLEIC ACID SEQUENCES CODING FOR SAID PEPTIDES
: TITLE OF INVENTION: PREPARATION AND UTILIZATION
: NUMBER OF SEQUENCES: 12
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Rhone-Poulenc Rorer Inc.
: STREET: 500 Arcola Road, 3C43
: CITY: Collegeville
: STATE: PA
: COUNTRY: USA
: ZIP: 19426
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: Macintosh
: OPERATING SYSTEM: System 7.1
: SOFTWARE: Word 5.1 (EPO PatentIn)
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/318,831
: FILING DATE: 19 October 1994
: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: FR92/04827
: FILING DATE: 21-Apr-1992
: ATTORNEY/AGENT INFORMATION:
: NAME: Smith, Julie K.
: REGISTRATION NUMBER: P-38,619
: REFERENCE/DOCKET NUMBER: ST92033-US
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (610) 454-3839
: TELEFAX: (610) 454-3808
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 2652 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: HYPOTHETICAL: NO
: ANTI-SENSE: NO
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 1..2445

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:      FEATURE:
:      NAME/KEY: CDS
:      LOCATION: 445..2445 (SEQ ID NO 3)
:
:      FEATURE:
:      NAME/KEY: CDS
:      LOCATION: 976..2445 (SEQ ID NO 4)
:
DS-08-318-831-1

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Query Match	9.9%	Score 36.6;	DB 1;	Length 2652;
Best Local Similarity	53.1%	Pred. No. 0.022;		
Matches 78;	Conservative 0;	Mismatches 69;	Indels 0;	Gaps 0;

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QY	TTTATTCTATATATGACATTTGGCGCGGAAACAAACAGCTGACCAAGAGCCCTCCAA	114
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QY	GCCAAATGTATAGTGTGAGACTCTCTACTATGTCATCCGTACTCTATGTATGATTTGA	174
Db	GACCCAGCTCTCTCCGGGAGCAATGTCTAGATATACCTATACATATCTGTTGGTTTCA	2521
QY	ATGTTCTCTTTCAATCATGCGCCATC	201
Db	TGGATTTCCTCTTCAGTATGTCTCTC	2548

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1  APPLICANT: Corley, Nell C.
2  TITLE OF INVENTION: DELAYED RECTIFIER POTASSIUM
3  TITLE OF INVENTION: CHANNEL HOMOLOG
4  NUMBER OF SEQUENCES: 4
5  CORRESPONDENCE ADDRESS:
6  ADDRESSEE: Incyte Pharmaceuticals, Inc.
7  STREET: 3174 Porter Drive
8  CITY: Palo Alto
9  STATE: CA
10 COUNTRY: USA
11 ZIP: 94304
12 COMPUTER READABLE FORM:
13 MEDIUM TYPE: Diskette
14 COMPUTER: IBM Compatible
15 OPERATING SYSTEM: DOS
16 SOFTWARE: FASTSEQ for Windows Version 2.0
17 CURRENT APPLICATION DATA:
18 APPLICATION NUMBER: US/09/069,896
19 FILING DATE:
20 CLASSIFICATION:
21 PRIOR APPLICATION DATA:
22 APPLICATION NUMBER:
23 FILING DATE:
24 ATTORNEY/AGENT INFORMATION:
25 NAME: Cerrone, Michael C
26 REGISTRATION NUMBER: 39,132
27 REFERENCE/DOCKET NUMBER: PP-0507 US
28 TELECOMMUNICATION INFORMATION:
29 TELEPHONE: 650-855-0555
30 TELEFAX: 650-845-4166
31 TELEX:
32 INFORMATION FOR SEQ ID NO: 2:
33 SEQUENCE CHARACTERISTICS:
34 LENGTH: 645 base pairs
35 TYPE: nucleic acid
36 STRANDEDNESS: single
37 TOPOLOGY: linear
38 IMMEDIATE SOURCE:
39 LIBRARY: BRSTNOT03
40 CLONE: 637471
41 US-09-069-896-2

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